

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 04:54:54 ; Search time 2762.24 Seconds
(without alignments)
65.021 Million cell updates/sec

Title: US-09-142-095-4

Perfect score: 19
Sequence: 1 ttgtctctgcagaggtt 19

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1028115 segs, 4726426750 residues

Total number of hits satisfying chosen parameters: 20456230

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: gb_est1:*
2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
5: gb_est5:*
6: gb_est6:*
7: gb_est7:*
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9: gb_est9:*
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257: gb_est158:*
258: gb_est159:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Genome Res. 6 (9), 791-806 (1996)
07044473

Contact: Soares, MB
Program for Rat Gene Discovery and Mapping
National Institutes of Health

451 Eckstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565

email: mscarets@bluewin.ch
 cDNA Library Preparation: M.B. Soares Lab Clone Distribution:
 clones will be available through Research Genetics (www.regen.com)
 This clone is also available through the I.M.A.G.E. Consortium at
 LLNL (info@image.llnl.gov). IMAGE ID- 1769283
 Seq primer: M13 Forward.

1. .522

```

/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/db_xref="taxon:10116"
/clone="UI-R-C2p-qt-e-07-0-UI"
/clone_1lb="UI-R-C2p"
/dev_stage="adult"

```

c	18	15.8	83.2	165	163	BE142610
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c	20	15.8	83.2	163	161	BE574964
c	21	15.8	83.2	251	161	BE580263
c	22	15.8	83.2	264	112	AM159504
c	23	15.8	83.2	287	116	BE488261
c	24	15.8	83.2	287	128	BE418963
c	25	15.8	83.2	290	134	BB466184
c	26	15.8	83.2	332	118	AM583615
c	27	15.8	83.2	352	169	BF808228
c	28	15.8	83.2	388	6	AA3378262
c	29	15.8	83.2	398	23	AI639055
c	30	15.8	83.2	401	170	BF857355
c	31	15.8	83.2	417	5	AA309827
c	32	15.8	83.2	428	190	W38798
c	33	15.8	83.2	440	175	BG334717
c	34	15.8	83.2	455	244	AA477763
c	35	15.8	83.2	464	228	AA043661
c	36	15.8	83.2	481	1	AA036818
c	37	15.8	83.2	487	159	NA4389
c	38	15.8	83.2	494	169	BF143356
c	39	15.8	83.2	495	24	AI172856
c	40	15.8	83.2	502	229	AA084899
c	41	15.8	83.2	504	105	AL049145
c	42	15.8	83.2	515	111	AM078344
c	43	15.8	83.2	521	233	AO172645
c	44	15.8	83.2	525	235	AO177005
c	45	15.8	83.2	549	230	AO533974
BE142610 IL0-H701						
AM306100 f179h04.y						
BB574964 BB574964						
BB580263 BB580263						
AM159504 za63c09.x						
AM482661 46483 MA1						
BB418963 BB218963						
BB466184 BB466184						
AM583615 l02a02.y						
BF802658 CMO-C7013						
AA3378262 ES909949						
AI639055 fX041265						
BF857435 OV3-F7021						
AA309827 ES1180840						
W38798 zbd7608.r1						
BG334717 dbb63B10.						
AA477763 LM029J5223						
AA456361 HS_5060.B						
AA036818 xK29a07.r						
NA4389 Y09a02.r1						
BF133356 IL2-BF086						
AI172856 fc3ze10.y						
AA084899 RPO1-11-2						
AL049145 DKP2P34F4F						
AM078344 fe48c07.y						
AO172645 HS_5566.A						
AO177005 nbd0067.L						
AO533974 RPO1-11-3						

156 a 117 c 104 g 145 t

Query Match	91.68;	Score 17.4;	DB 150;	Length 522;
Best Local Similarity	94.78;	Pred. No. 2.1e+02;		
Matches 18; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0;

```
Dy      1 ttgtctcctgccagagtt   19  
         ||| | | | | | | | |  
Db     128 TTGGCTCCTGCCAGAGGT   146
```

RESULT	1
BF545097	
LOCUS	BF545097 522 bp mRNA
DEFINITION	U1-R-C2p-qt-e-07-0-01 r1 U1-R-C2p Rattus norvegicus cDNA clone
ACCESSION	U1-R-C2p-qt-e-07-0-01 5', mRNA sequence.
VERSION	BF545097
KEYWORDS	BF545097.1 GI:11636204
SOURCE	EST.
ORGANISM	Norway rat.
	Rattus norvegicus
	11-DEC-2000

REFERENCE: 1. (b) (8) 1 to 5221

AUTHORS	TITLE
Bonafide, M.F., Lennon, G. and Soares, M.B.	Normalisation and subtraction: two approaches to facilitate gene

RESULT	2
LOCUS	AO540572/c
DEFINITION	AO540572 608 bp DNA
ACCESSION	RFCT-11-358E23.TV RFCT-11 Homo sapiens genomic clone RFCT-11-358E23
VERSION	AO540572
KEYWORDS	AO540572.1 GI:4871102
SOURCE	GSS.
ORGANISM	human.
	Homo sapiens

REFERENCE
AUTHORS
1 (bases 1 to 608)
Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter

TITLE
JOURNAL
COMMENT

J.C.
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building (1997)
Unpublished (1997)
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: dbeetlgr.org

Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
Research Genet cs (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.
Seq primer: 17
Class: BAC ends.

FEATURES
source

1. 608
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="GDB:7637206"
/db_xref="taxon:9606"
/clone="RPCI-11-358E23"
/clone_11b="RPCI-11"
/sex="Male"
/cell_type="lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"
BASE COUNT 259 a 83 c 93 g 173 t
ORIGIN

Query Match 91.6%; Score 17.4; DB 230; Length 608;
Best Local Similarity 94.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 1 ttgtctctgcagaggtt 19
|||||
Db 138 TTGGCTCTCTCCAGAGTT 120

RESULT 3
AW429811 367 bp mRNA EST 09-JUL-2000
LOCUS 68306 MARC 1P1G Sus scrofa cDNA 5', mRNA sequence.
DEFINITION
ACCESSION AW429811
VERSION
KEYWORDS
SOURCE
ORGANISM

Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.
Fahnenkrug, S.C., Fekking, B.A., Rohrer, G.A., Smith, T.P.L., Casas, E.,
Stone, R.T., Heaton, M.P., Grose, W.M., Bennett, G.A., Laegreid, W.W.
and Keeler, J.W.
Design and use of two pooled tissue normalized cDNA libraries for
EST discovery in swine
Unpublished (2000)
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@mail.marc.usda.gov
Single pass sequencing. Bases called and trimmed with phred
v0.980904.e. Vector identified by cross-match with the -m1nscore 20
PCR Primers
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACAT
BACKWARD: GTTTCACAGTCAGCAGC

FEATURES
source

Plate: 28 row: C column: 15
Seq primer: ATTAGCTGACACTATAG.
Location/Qualifiers
1. 367
/organism="Sus scrofa"
/db_xref="taxon:9823"
/clone_11b="MARC 1P1G"
/tissue_type="pooled"
/lab_host="DH10B"
/note="Vector: PCMV SPOR6; Site_1: XbaI; Site_2: XhoI;
Library made from pooled tissue from day 11, 13, 15, 20,
and 30 embryos."
BASE COUNT 91 a 100 c 115 g 60 t 1 others
ORIGIN

Query Match 89.5%; Score 17; DB 115; Length 367;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 tgcctctgcagaggtt 19
|||||
Db 256 TGCTCTCTCCAGAGTT 272

RESULT 4
BG297415 936 bp mRNA EST 21-FEB-2001
LOCUS 60295559P1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4507022 5',
DEFINITION
ACCESSION BG297415
VERSION
KEYWORDS
SOURCE
ORGANISM

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: The Cepko Laboratory
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
Plate: LHAM0383 row: f column: 15
High quality sequence stop: 604.
Location/Qualifiers
1. 936
/organism="Mus musculus"
/db_xref="taxon:10090"
/clone="IMAGE:4507022"
/clone_11b="NIH_MGC_94"
/tissue_type="retina"
/lab_host="DH10B (phage-resistant)"
/note="Organ: eye; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 3.3 kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH_MGC Library."

FEATURES
source

Query Match 89.5%; Score 17; DB 175; Length 936;
Best Local Similarity 100.0%; Pred. No. 3.6e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
BASE COUNT 367 a 218 c 259 g 92 t
ORIGIN

PT Improving drug trial efficiency comprises identifying participants

ps Examples; Page 27-29; 63pp; English.

PE
XX
10-JAN-1992; 92MO-050028

PR 10-JAN-1991; 91US-0639453.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICE.
 XX
 PI Owens IS, Ritter JK;
 XX
 DR WPI: 1992-284593/34.
 DR P-PSDB; AAR30194.
 XX
 PT Isolated gene locus UGT1, DNA segments and diagnostic probes -
 PT for diagnosing Gilbert's disease and Crigler-Najjar syndrome
 PT types I and II
 XX
 PS Disclosure: Fig 1F; 99pp; English.
 XX
 CC The isolated gene locus, UGT1, has a sequence of about 10000 bp
 CC which represent (1) Exon 1, comprising 6 transcriptional units
 CC (UGT1F, E, D, C, BP and A), represented in AA027368 and
 CC AA033020-24 respectively;
 CC (2) Exon 2, represented in AA033025;
 CC (3) Exon 3, represented in AA033026;
 CC (4) Exon 4, represented in AA033026;
 CC (5) Exon 5, represented in AA033027; and
 CC (6) about 69 kb of non-sequenced DNA.
 CC Six unique N-terminal of 286-289 amino acids are encoded by
 CC the six different first exons and identical C-terminal of 246 amino
 CC acids are encoded by the common exons 2-5. The UGT1 gene locus
 CC encodes a family of UDP-glucuronosyl transferase isozymes, two of
 CC which metabolise bilirubin.
 CC Patients having Crigler-Najjar Syndrome (CN) Type I, have a
 CC mutation present in the second common exon.
 CC
 SQ Sequence 1167 BP; 255 A; 259 C; 272 G; 340 T; 41 other;
 XX
 XX

Query Match 100.0%; Score 21; DB 13; Length 1167;
 Best Local Similarity 100.0%; Pred. No. 0.19; 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0;

QY 1 cccactggagatcaacagatct 21
 Db 286 CCACCTGGGATCAACAGATCT 266.

RESULT 4
 AA027369/C
 ID AA027369 standard; cDNA; 2351 BP.
 XX
 AC AA027369;
 XX
 DT 27-JAN-1993 (first entry)
 XX
 DE HUG-Brl.
 XX
 KM Bilirubin: UDP-glucuronosyltransferase; HUGBrl; HUGB2;
 KM monoglucuronide; diglucuronide; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FH CDS 16..784
 FT CDS
 FT polyA_signal 2330..2335
 FT /tag- a
 FT /number- b
 FT polyA_signal 2338..2343
 FT /tag- c
 FT /number- 2
 XX
 XX
 PN MO9212987-A.
 XX
 PD 06-AUG-1992.

PF 10-JAN-1992; 92MO-US00282.
 XX
 PR 10-JAN-1991; 91US-0639453.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICE.
 XX
 PI Owens IS, Ritter JK;
 XX
 DR WPI: 1992-284593/34.
 DR P-PSDB; AAR26153.
 XX
 PT Isolated gene locus UGT1, DNA segments and diagnostic probes -
 PT for diagnosing Gilbert's disease and Crigler-Najjar syndrome
 PT types I and II
 XX
 PS Disclosure: Fig 9A-I; 99pp; English.
 XX
 CC Two human liver bilirubin UDP-glucuronosyltransferase cDNAs have
 CC been isolated. They are referred to as HUGBrl (AA027369) and HUGB2
 CC (AA027370) (Ritter, et al., J. Biol. Chem. 266:1043-1047 (1991)) and,
 CC upon expression individually in COS-1 cells, encode isoforms that
 CC catalyse the formation of the two bilirubin monoglucuronides and
 CC the diglucuronide.
 CC The cDNAs contain identical 3' ends (1469 bp in length) to each
 CC other and to that of the human phenol transferase cDNA, HUGP1
 CC (Harding et al., Proc. Natl. Acad. Sci. USA 85:8281 (1988)).
 CC In contrast, they have unique 5' ends.
 CC
 SQ Sequence 2351 BP; 602 A; 540 C; 556 G; 653 T; 0 other;
 XX
 XX

Query Match 100.0%; Score 21; DB 13; Length 2351;
 Best Local Similarity 100.0%; Pred. No. 0.21;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 cccactggagatcaacagatct 21
 Db 121 CCACCTGGGATCAACAGATCT 101

RESULT 5
 AAD00564
 ID AAD00564 standard; DNA; 504 BP.
 XX
 AC AAD00564;
 XX
 DT 29-AUG-2000 (first entry)
 XX
 DE Streptococcus pyogenes strain KTL3 partial GRAB protein encoding DNA.
 XX
 KM GRAB protein; protein G related alpha2m binding protein; vaccine;
 KM alpha2-macroglobulin; group A Streptococcus; GAS; antibiotic;
 KM Immune response; Streptococcus pyogenes infection; ds.
 XX
 OS Streptococcus pyogenes.
 XX
 FH Key Location/Qualifiers
 FH CDS 1..504
 FT CDS
 FT /tag- a
 FT /product- "GRAB protein"
 FT /partial
 XX
 PN MO200026240-A2.
 XX
 PD 11-MAY-2000.
 XX
 PD 02-NOV-1999; 99MO-GB03631.
 XX
 PD 02-NOV-1998; 98GB-0023975.
 XX
 PA (ACTI-) ACTINOVA LTD.
 XX
 PI Bjorck LH, Rasmussen M;

XX NPI; 2000-365572/31.
 DR P-SDB; AAV71046.
 PT New alpha2M binding protein for generating a protective immune response
 PT to group A streptococcus and purifying the binding protein
 XX
 PS Claim 13; Page 65; 67pp; English.
 XX
 CC The patent discloses a new family of proteins termed GRAB (protein G
 CC related alpha2M binding protein) from Streptococcus pyogenes which have
 CC the ability to bind alpha2-macroglobulin (alpha2M) and show homology to
 CC protein G of group G Streptococcus. GRAB protein and peptides derived
 CC from it are used in vaccine compositions for generating a protective
 CC immune response against group A Streptococcus. Antibodies against GRAB
 CC are useful for treating Streptococcus pyogenes infections. The protein
 CC is also useful for purifying alpha2M from a sample. The present sequence
 CC is a DNA encoding partial GRAB protein from S. pyogenes strain KTL3.
 CC The protein has alpha2M binding region and is useful in vaccine
 CC composition.
 CC
 XX
 SQ Sequence 504 BP; 188 A; 97 C; 108 G; 111 T; 0 other;

Query Match 75.2%; Score 15.8; DB 21; Length 504;
 Best Local Similarity 89.5%; Pred. No. 74;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggagcacacagatctc 21
 | ||||| ||||| |||||
 Db 12 agtgggttcacacagatctc 30

RESULT 6
 AAX30352
 ID AAX30352 standard; DNA; 697 BP.
 XX
 AC AAX30352;
 XX
 DT 14-MAY-1999 (first entry)
 XX
 DE DNA encoding a human secreted protein.
 XX
 KW Secreted protein; cancer; tumour; neurodegenerative disorder;
 KW developmental abnormality; foetal deficiency; blood disorder;
 KW CNS disorder; immune system disease; autoimmune disease; hepatic disease;
 KW renal disease; diabetes; inflammation; allergy; ischemic shock;
 KW Alzheimer's; cognitive disorder; schizophrenia; cardiovascular disorder;
 KW prostate disease; asthma; osteoporosis; arthritis; ss.
 KW
 OS Homo sapiens.
 XX
 FN WO9907881-A1.
 XX
 PD 18-FEB-1999.
 PD
 XX
 PF 04-AUG-1998; 98MO-US16235.
 XX
 PR 19-AUG-1997; 97US-0056732.
 PR 05-AUG-1997; 97US-0054798.
 PR 05-AUG-1997; 97US-0054803.
 PR 05-AUG-1997; 97US-0054804.
 PR 05-AUG-1997; 97US-0054806.
 PR 05-AUG-1997; 97US-0054807.
 PR 05-AUG-1997; 97US-0054808.
 PR 05-AUG-1997; 97US-0054809.
 PR 05-AUG-1997; 97US-0055309.
 PR 05-AUG-1997; 97US-0055310.
 PR 05-AUG-1997; 97US-0055312.
 PR 05-AUG-1997; 97US-0055386.
 PR 05-AUG-1997; 97US-0055311.
 PR 18-AUG-1997; 97US-0055970.
 PR 18-AUG-1997; 97US-0055986.

PR 19-AUG-1997; 97US-0056365.
 PR 19-AUG-1997; 97US-0056366.
 PR 19-AUG-1997; 97US-0056557.
 PR 19-AUG-1997; 97US-0056370.
 PR 19-AUG-1997; 97US-0056371.
 PR 19-AUG-1997; 97US-0056563.
 PR 19-AUG-1997; 97US-0056731.
 XX
 PA (HDMA-) HUMAN GENOME SCI INC.
 XX
 PI Brewer LA, Ebner R, Ferlie AM, Greene JM, Janat F, Ni J;
 PI Olsen HS, Rosen CA, Ruben SM, Soppet DR, Young PE, Yu G;
 DR NPI; 1999-167452/14.
 DR P-SDB; AAV10831.
 XX
 PT New isolated human genes encoding secreted polypeptides - useful for
 PT diagnosis and treatment of pathological diseases
 XX
 PS Claim 3; Page 247; 331pp; English.
 XX
 CC The specification describes secreted proteins and their corresponding
 CC polynucleotides which are useful for preventing, treating or ameliorating
 CC medical conditions, e.g. by protein or gene therapy. Pathological
 CC conditions can also be diagnosed by determining the amount of the
 CC secreted polypeptides in a sample or by determining the presence of
 CC mutations in the polynucleotides. Specific uses are described for each
 CC of the products, based on which tissues they are most highly
 CC expressed in, and include developing products for the diagnosis or
 CC treatment of cancer, tumours, neurodegenerative disorders, developmental
 CC abnormalities and foetal deficiencies, blood disorders, CNS disorders,
 CC diseases of the immune system, autoimmune diseases, hepatic and renal
 CC disease, diabetes, inflammation, allergies, ischemic shock, Alzheimer's
 CC and cognitive disorders, schizophrenia, cardiovascular disorders,
 CC prostate diseases, asthma, disorders involving osteoclasts such as
 CC osteoporosis, arthritis or malignancies, diseases of testes, lung or
 CC thymus, digestive/endocrine disorders, infections and AIDS. The
 CC polypeptides are also useful for identifying their binding partners.
 CC
 XX
 SQ Sequence 697 BP; 233 A; 121 C; 171 G; 169 T; 3 other;

Query Match 75.2%; Score 15.8; DB 20; Length 697;
 Best Local Similarity 89.5%; Pred. No. 77;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 cactggagcacacagatc 20
 ||| ||||| ||||| |||||
 Db 274 cactggagcacacagatc 292

RESULT 7
 AAV30310
 ID AAV30310 standard; DNA; 1073 BP.
 XX
 AC AAV30310;
 XX
 DT 28-SEP-1998 (first entry)
 XX
 DE Bacillus thuringiensis MIS-6 insecticidal toxin 196F3 DNA.
 XX
 KW Insecticide; pesticide; toxin; MIS-6; delta-endotoxin;
 KW biological control; lepidopteran; coleopteran; ss.
 XX
 OS Bacillus thuringiensis strain PS196F3 (NRRL B-21872).
 XX
 XX
 FT Key Location/Qualifiers
 FT CDS 2..1072
 XX /*tag- a
 PN WO9818932-A2.
 XX
 PD 07-MAY-1998.

XX 30-OCT-1997; 97WO-US19804.
 XX 30-OCT-1996; 96US-0029848.
 PR (MICO) MYCOGEN CORP.
 XX
 XX Dullum CJ, Feltelson JS, Loewer D, Muller-Cohn J;
 PI Narva KE, Schmelts JL, Schnepf HE, Schwab G, Stamp L;
 PI Stockhoff BA;
 XX
 DR WPI: 1998-272226/24.
 DR P-PSDB; AAW60227.
 XX
 PT Bacillus thuringiensis isolates - used for producing pesticidal
 PT toxins and nucleotide sequences for control of lepidopterans and
 PT coleopterans
 XX
 PS Claim 5; Page 90; 139pp; English.
 XX
 CC This DNA sequence encodes a novel soluble toxin (see AAW60227) of
 CC Bacillus thuringiensis (B.t.) strain PS196F3 (NRRL B-21872). This
 CC toxin belongs to the novel MIS-6 family of B.t. toxins that have
 CC toxicity against non-mammalian pests. The novel DNA was obtained
 CC by PCR amplification (see AAV30298-99) of total cellular genomic
 CC DNA. It can be used to produce recombinant hosts (preferably plant
 CC or bacterial) that express the toxin, or as a PCR primer or
 CC hybridisation probe for use in identifying and characterising
 CC MIS-6 family toxin genes. Disclosed and claimed are novel B.t.
 CC isolates and toxins (see AAW60218-32) which have activity against
 CC lepidopteran and/or coleopteran pests, isolated genes, probes
 CC and primers (see AAV30288-321 and AAV9734-87), and transformed host
 CC cells. The invention provides 8 entirely new families of toxins,
 CC including MIS-6, from B.t. isolates. MIS toxins are also useful
 CC for their ability to form pores in cell membranes, and can be used
 CC to facilitate entry of a second agent into a target cell.
 CC
 SO Sequence 1073 BP; 380 A; 201 C; 215 G; 270 T; 7 other;

Query Match 75.2%; Score 15.8; DB 19; Length 1073;
 Best Local Similarity 89.5%; Pred. No. 82;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 actggatcacagatct 21
 |||||
 Db 838 actggatcacagatct 856

RESULT 8
 AAV70397
 ID AAV70397 standard; cDNA; 4351 BP.
 XX
 AC AAV70397;
 XX
 DT 10-FEB-1999 (first entry)
 XX
 DE LRP5 isoform 2 longest open reading frame (also isoform 4,5,6).
 XX
 XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;
 KW endocytosis; insulin dependent diabetes mellitus; autoimmune disease;
 KW glomerulonephritis; inflammation; viral infection; osteoporosis;
 KW hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.
 XX
 OS Homo sapiens.
 XX
 PN MO9846743-A1.
 XX
 PD 22-OCT-1998.
 XX
 PR 15-APR-1998; 98WO-GB01102.
 XX
 PR 05-JUN-1997; 97US-0048740.
 XX

PR 15-APR-1997; 97US-0043553.
 XX
 PA (MERI) MERCK & CO INC.
 PA (WEIL) WEILCOME TRUST LTD.
 XX
 PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JM;
 PI Hey P, Kawaguchi Y, Merriam TR, Metzker ML, Nakagawa Y;
 PI Phillips MS, Todd JA, Twells RCU;
 XX
 DR WPI: 1998-594573/50.
 XX
 PT New isolated LDL-receptor related protein - used to develop products
 PT for treating e.g. elevated triglyceride levels, diabetes,
 PT autoimmune disorders, inflammation or Alzheimer's disease
 XX
 PS Claim 7; Fig 11b; 200pp; English.
 XX
 CC The present sequence represents LRP5 (low density lipoprotein (LDL)
 CC receptor related protein, previously designated LRP-3) isoform 2 cDNA.
 CC Nucleic acid molecules (NAMS) encoding LRP5 can be used for determining
 CC if an individual is susceptible to insulin dependent diabetes mellitus
 CC (IDDM). The NAMS or proteins can be used for reducing triglyceride levels
 CC in the serum of an individual. Therapies that affect LRP5 may also be
 CC useful in the treatment of autoimmune diseases such as
 CC glomerulonephritis, diseases and disorders involving disruption of
 CC endocytosis and/or antigen presentation, cytokine clearance and/or
 CC inflammation, viral infection, pathogenic bacterial toxin contamination,
 CC elevation of free fatty acids or hypercholesterolemia, type 2 diabetes,
 CC osteoporosis, Alzheimer's disease and cardiovascular disease. Products
 CC from the present invention can also be used for detection, diagnosis and
 CC drug screening.
 CC
 SO Sequence 4351 BP; 875 A; 1435 C; 1309 G; 732 T; 0 other;

Query Match 75.2%; Score 15.8; DB 19; Length 4351;
 Best Local Similarity 89.5%; Pred. No. 1e+02;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 cactggatcacagatc 20
 |||||
 Db 1716 cactggatcacagatc 1734

RESULT 9
 AAV70395
 ID AAV70395 standard; cDNA; 4843 BP.
 XX
 AC AAV70395;
 XX
 DT 10-FEB-1999 (first entry)
 XX
 DE LRP5 cDNA longest open reading frame.
 XX
 XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;
 KW endocytosis; insulin dependent diabetes mellitus; autoimmune disease;
 KW glomerulonephritis; inflammation; viral infection; osteoporosis;
 KW hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.
 XX
 OS Homo sapiens.
 XX
 PN MO9846743-A1.
 XX
 PD 22-OCT-1998.
 XX
 PR 15-APR-1998; 98WO-GB01102.
 XX
 PR 05-JUN-1997; 97US-0048740.
 XX
 PR 15-APR-1997; 97US-0043553.
 XX
 PA (MERI) MERCK & CO INC.
 PA (WEIL) WEILCOME TRUST LTD.
 XX

PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JM;
 PI Hey P, Kawaguchi Y, Merriman TR, Metzker ML, Nakagawa Y;
 PI Phillips MS, Todd JA, Twells RCJ;
 XX
 DR WPI: 1998-594573/50.

XX
 PT New isolated LDL-receptor related protein - used to develop products
 PT for treating, e.g. elevated triglyceride levels, diabetes,
 PT autoimmune disorders, inflammation or Alzheimer's disease
 XX
 PS Example 1: Fig 5b: 200pp; English.

XX The present sequence represents the longest open reading frame of LRP5
 CC (low density lipoprotein (LDL) receptor related protein, previously
 CC designated LRP-3) CDNA from the present invention. Nucleic acid
 CC molecules (NMs) encoding LRP5 can be used for determining if an
 CC individual is susceptible to insulin dependent diabetes mellitus (IDDM).
 CC The NMs or proteins can be used for reducing triglyceride levels
 CC in the serum of an individual. Therapies that affect LRP5 may also be
 CC useful in the treatment of autoimmune diseases such as
 CC glomerulonephritis, diseases and disorders involving disruption of
 CC endocytosis and/or antigen presentation, cytokine clearance and/or
 CC inflammation, viral infection, pathogenic bacterial toxin contamination,
 CC elevation of free fatty acids or hypercholesterolemia, type 2 diabetes,
 CC osteoporosis, Alzheimer's disease and cardiovascular disease. Products
 CC from the present invention can also be used for detection, diagnosis and
 CC drug screening.

XX Sequence 4843 BP; 953 A; 1601 C; 1478 G; 811 T; 0 other;

Query Match 75.2%; Score 15.8; DB 19; Length 4843;

Best Local Similarity 89.5%; Pred. No. 1e+02; 2; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 cactgggacacagatc 20

Db 2207 cactgggacacagatc 2225

RESULT 10

AAV70398 standard; CDNA; 4915 BP.

AC AAV70398;

DT 10-FEB-1999 (first entry)

DE LRP5 isoform 3 putative open reading frame.

XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;

KW endocytosis; insulin dependent diabetes mellitus; autoimmune disease;

KW glomerulonephritis; inflammation; viral infection; osteoporosis;

KW hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.

OS Homo sapiens.

PN WO9846743-A1.

PD 22-OCT-1998.

PF 15-APR-1998; 98WO-GB01102.

PR 05-JUN-1997; 97US-0048740.

PA 15-APR-1997; 97US-0043553.

PA (MERI) MERCK & CO INC.

PA (WELL) WELLCOME TRUST LTD.

PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JM;

PI Hey P, Kawaguchi Y, Merriman TR, Metzker ML, Nakagawa Y;

PI Phillips MS, Todd JA, Twells RCJ;

DR WPI: 1998-594573/50.
 XX
 PT New isolated LDL-receptor related protein - used to develop products
 PT for treating, e.g. elevated triglyceride levels, diabetes,
 PT autoimmune disorders, inflammation or Alzheimer's disease
 XX
 PS Example 1: Fig 12c: 200pp; English.

XX The present invention describes LRP5 (low density lipoprotein (LDL)
 CC receptor related protein, previously designated LRP-3). The present
 CC sequence represents the putative open reading frame of LRP5 isoform 3.
 CC Nucleic acid molecules (NMs) encoding LRP5 can be used for determining
 CC if an individual is susceptible to insulin dependent diabetes mellitus
 CC (IDDM). The NMs or proteins can be used for reducing triglyceride levels
 CC in the serum of an individual. Therapies that affect LRP5 may also be
 CC useful in the treatment of autoimmune diseases such as
 CC glomerulonephritis, diseases and disorders involving disruption of
 CC endocytosis and/or antigen presentation, cytokine clearance and/or
 CC inflammation, viral infection, pathogenic bacterial toxin contamination,
 CC elevation of free fatty acids or hypercholesterolemia, type 2 diabetes,
 CC osteoporosis, Alzheimer's disease and cardiovascular disease. Products
 CC from the present invention can also be used for detection, diagnosis and
 CC drug screening.

SO Sequence 4915 BP; 992 A; 1614 C; 1474 G; 835 T; 0 other;

Query Match 75.2%; Score 15.8; DB 19; Length 4915;

Best Local Similarity 89.5%; Pred. No. 1e+02; 2; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 cactgggacacagatc 20

Db 2279 cactgggacacagatc 2297

RESULT 11

AAV85549 standard; CDNA; 5022 BP.

AC AAV85549;

DT 10-FEB-1999 (first entry)

DE LRP5 isoform 5 CDNA.

XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;

KW endocytosis; insulin dependent diabetes mellitus; autoimmune disease;

KW glomerulonephritis; inflammation; viral infection; osteoporosis;

KW hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.

OS Homo sapiens.

PN WO9846743-A1.

PD 22-OCT-1998.

PF 15-APR-1998; 98WO-GB01102.

PR 05-JUN-1997; 97US-0048740.

PA 15-APR-1997; 97US-0043553.

PA (MERI) MERCK & CO INC.

PA (WELL) WELLCOME TRUST LTD.

PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JM;

PI Hey P, Kawaguchi Y, Merriman TR, Metzker ML, Nakagawa Y;

PI Phillips MS, Todd JA, Twells RCJ;

DR WPI: 1998-594573/50.

PT New isolated LDL-receptor related protein - used to develop products
 PT for treating, e.g. elevated triglyceride levels, diabetes,

PT autoimmune disorders, inflammation or Alzheimer's disease
XX
PS Claim 7; Fig 14; 200pp; English.

CC The present invention describes LRP5 (low density lipoprotein (LDL)
CC receptor related protein, previously designated LRP-3). The present
CC sequence represents the LRP5 isoform 5 cDNA.
CC Nucleic acid molecules (NMs) encoding LRP5 may also be
CC if an individual is susceptible to insulin dependent diabetes mellitus
CC (IDDM). The NMs or proteins can be used for reducing triglyceride levels
CC in the serum of an individual. Therapies that affect LRP5 may also be
CC useful in the treatment of autoimmune diseases such as
CC glomerulonephritis, diseases and disorders involving disruption of
CC endocytosis and/or antigen presentation, cytokine clearance and/or
CC inflammation, viral infection, pathogenic bacterial toxin contamination,
CC elevation of free fatty acids or hypercholesterolemia, type 2 diabetes,
CC osteoporosis, Alzheimer's disease and cardiovascular disease. Products
CC from the present invention can also be used for detection, diagnosis and
CC drug screening.

SO Sequence 5022 BP; 1036 A; 1606 C; 1503 G; 877 T; 0 other;

Query Match 75.2%; Score 15.8; DB 19; Length 5022;
Best Local Similarity 89.5%; Pred. No. 1e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 cactgggacacagatc 20
||||| ||||| |||
DB 2178 cactgggacacagatc 2196

RESULT 12

AAV70396
ID AAV70396 standard; cDNA; 5098 BP.

XX AAV70396;

XX 10-FEB-1999 (first entry)

DE LRP5 isoform 1 cDNA.

XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;
KM endocytosis; insulin dependent diabetes mellitus; autoimmune disease;
KM glomerulonephritis; inflammation; viral infection; osteoporosis;
KM hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.

OS Homo sapiens.

PN W09846743-A1.

PD 22-OCT-1998.

PF 15-APR-1998; 98WO-GB01102.

PR 05-JUN-1997; 97US-0048740.

PR 15-APR-1997; 97US-0043553.

PA (MERI) MERCK & CO INC.

PI (WELL) WELLCOME TRUST LTD.

PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JW;

PI Hey P, Kawaguchi Y, Merriman TR, Metzker ML, Nakagawa Y;

PI Phillips MS, Todd JA, Twells RCJ;

XX WPI; 1998-594573/50.

PS New isolated LDL-receptor related protein - used to develop products
XX for treating, e.g. elevated triglyceride levels, diabetes,
XX autoimmune disorders, inflammation or Alzheimer's disease
XX Claim 3; Fig 5a; 200pp; English.

CC The present sequence represents LRP5 (low density lipoprotein (LDL)
CC receptor related protein, previously designated LRP-3) isoform 1 cDNA.
CC Nucleic acid molecules (NMs) encoding LRP5 can be used for determining
CC if an individual is susceptible to insulin dependent diabetes mellitus
CC (IDDM). The NMs or proteins can be used for reducing triglyceride levels
CC in the serum of an individual. Therapies that affect LRP5 may also be
CC useful in the treatment of autoimmune diseases such as
CC glomerulonephritis, diseases and disorders involving disruption of
CC endocytosis and/or antigen presentation, cytokine clearance and/or
CC inflammation, viral infection, pathogenic bacterial toxin contamination,
CC elevation of free fatty acids or hypercholesterolemia, type 2 diabetes,
CC osteoporosis, Alzheimer's disease and cardiovascular disease. Products
CC from the present invention can also be used for detection, diagnosis and
CC drug screening.

SO Sequence 5098 BP; 1035 A; 1645 C; 1541 G; 877 T; 0 other;

Query Match 75.2%; Score 15.8; DB 19; Length 5098;
Best Local Similarity 89.5%; Pred. No. 1e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 cactgggacacagatc 20
||||| ||||| |||
DB 2255 cactgggacacagatc 2273

RESULT 13

AAV86019
ID AAV86019 standard; cDNA; 5125 BP.

XX AAV86019;

XX 10-FEB-1999 (first entry)

DE Lrp5 isoform 3 cDNA.

XX LRP5; LDL-receptor related protein; LRP-3; IDDM; diagnosis;
KM endocytosis; insulin dependent diabetes mellitus; autoimmune disease;
KM glomerulonephritis; inflammation; viral infection; osteoporosis;
KM hypercholesterolemia; Alzheimer's disease; low density lipoprotein; ss.

OS Homo sapiens.

PN W09846743-A1.

PD 22-OCT-1998.

PF 15-APR-1998; 98WO-GB01102.

PR 05-JUN-1997; 97US-0048740.

PR 15-APR-1997; 97US-0043553.

PA (MERI) MERCK & CO INC.

PI (WELL) WELLCOME TRUST LTD.

PI Caskey CT, Cox RD, Gerhold D, Hammond H, Hess JW;

PI Hey P, Kawaguchi Y, Merriman TR, Metzker ML, Nakagawa Y;

PI Phillips MS, Todd JA, Twells RCJ;

XX WPI; 1998-594573/50.

PS New isolated LDL-receptor related protein - used to develop products
XX for treating, e.g. elevated triglyceride levels, diabetes,
XX autoimmune disorders, inflammation or Alzheimer's disease
XX Example 1; Fig 12a; 200pp; English.

CC The present invention describes LRP5 (low density lipoprotein (LDL)
CC receptor related protein, previously designated LRP-3). The present
CC sequence represents the LRP5 isoform 3 cDNA.
CC Nucleic acid molecules (NMs) encoding LRP5 can be used for determining
CC if an individual is susceptible to insulin dependent diabetes mellitus

CC from the present invention can also be used for detection, diagnosis and
CC drug screening.

XX
SQ Sequence 5166 BP; 1082 A; 1654 C; 1530 G; 899 T; 1 other;

Query Match 75.2%; Score 15.8; DB 19; Length 5166;
Best Local Similarity 89.5%; Pred. No. 1e+02;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 cactgggacacagatc 20
|||||

Db 2333 cactgggacacagatc 2351

Search completed: July 25, 2001, 05:23:04
Job time: 4677 sec

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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:16:42; Search time 1290.33 Seconds

(without alignments)
251.736 Million cell updates/sec

Title: US-09-142-095-2

Perfect score: 21

Sequence: 1 ccactggatcacacatctct 21

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 1344157 segs, 7733874588 residues 2688314

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database:

GenEmb1:*
1: gb_ba1:*
2: gb_ba2:*
3: gb_ba3:*
4: gb_in1:*
5: gb_in2:*
6: gb_in3:*
7: gb_om:*
8: gb_ov:*
9: gb_pat1:*
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13: gb_pl2:*
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19: em_higo_hum:*
20: em_higo_inv:*
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37: em_hum4:*
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39: em_hum6:*
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41: em_in:*
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45: em_pat:*
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48: em_rod:*
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51: em_un:*
52: em_v1:*
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73: gb_hg14:*
74: gb_hg15:*
75: gb_hg16:*
76: gb_hg17:*
77: gb_hg18:*
78: gb_hg19:*
79: gb_hg20:*
80: gb_hg21:*
81: gb_hg22:*
82: gb_hg23:*
83: gb_hg24:*
84: gb_hg25:*
85: gb_pr1:*
86: gb_pr2:*
87: gb_pr3:*
88: gb_pr4:*
89: gb_pr5:*
90: gb_pr6:*
91: gb_pr7:*
92: gb_pr8:*
93: gb_pr9:*
94: gb_pr10:*
95: gb_rod:*
96: gb_in4:*
97: gb_pr10:*
98: em_ba3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	21	100.0	21	9 A65501	A65501 Sequence 2
2	21	100.0	292	89 AF357220	AF357220 Homo sapi
3	21	100.0	531	89 AF357295	AF357295 Homo sapi
4	21	100.0	541	88 AF180372	AF180372 Homo sapi
5	21	100.0	918	88 AF110194	AF110194 Homo sapi
6	21	100.0	1190	97 HOMOG71A	M84125 Human billir
7	21	100.0	2351	97 HOMOG71A	M57899 Human billir
8	21	100.0	3341	91 D87674	D87674 Homo sapien

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9 21 100.0 68770 86 AC006985
10 21 100.0 176619 70 AC026497
11 21 100.0 198872 89 AF297093
12 19.4 1602 91 AF029168
13 19.4 1647 88 AF104339
14 17.8 84.8 104480 60 AC007905
15 17.8 82.9 12887 93 HSD137K2
16 17.4 82.9 12887 1 AF077006
17 17 81.0 163245 63 AC013699
18 17 81.0 167230 79 AL355679
19 17 81.0 168656 90 AL359922
20 16.8 80.0 795 53 CDS0727X
21 16.8 80.0 818 71 AC035580
22 16.8 80.0 900 72 AC054151
23 16.8 80.0 999 53 CDS07102
24 16.8 80.0 1721 7 AF232676
25 15.8 80.0 17988 5 CEC3303
26 16.8 80.0 33513 63 AC012901
27 16.8 80.0 66168 6 DMSE30007
28 16.8 80.0 66857 76 AC054799_3
29 16.8 80.0 123926 76 AC079633
30 16.8 80.0 14870 92 HS173A13
31 16.8 80.0 151441 71 AC036158
32 16.8 80.0 161419 84 CDS0188X
33 16.8 80.0 162960 79 AL158062
34 16.8 80.0 163749 79 AL157833
35 16.8 80.0 173965 90 AL159977
36 16.8 80.0 182007 82 AL1590125
37 16.8 80.0 193260 69 AC025598
38 16.8 80.0 199648 72 AC060771
39 16.8 80.0 201530 83 CDS01DXK
40 16.8 80.0 294218 4 AE003430
41 16.4 78.1 6127 59 IBC0961
42 16.4 78.1 9081 58 AIBVCG
43 16.4 78.1 20500 59 IBACGB
44 16.4 78.1 27608 59 IBACGB
45 16.4 78.1 87960 65 AC017906

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ALIGNMENTS

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RESULT 1
LOCUS A65501 21 bp DNA
DEFINITION Sequence 2 from Patent WO9732042.
ACCESSION A65501
VERSION A65501.1 GI:4531236
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 21)
AUTHORS Burchell, B.
TITLE DRUG TRIAL ASSAY SYSTEM
JOURNAL Patent NO 9732042-A 2 04-SEP-1997;
UNIV DUNDEE (GB)
COMMENT Other publication AU 2224197 19970916.
FEATURES
source
1..21
/organism="unidentified"
/db_xref="taxon:33644"

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BASE COUNT 6 a 6 c 4 g 5 t
ORIGIN

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Query Match 100.0%; Score 21; DB 9; Length 21;
Best Local Similarity 100.0%; Pred. No. 0.68;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 ccaactggatcaacagatct 21
DB 1 ccaactggatcaacagatct 21

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```

RESULT 2
LOCUS AF357220/c
DEFINITION Homo sapiens bilirubin UDP-glucuronosyltransferase 1 (UGT1) gene,
ACCESSION AF357220
VERSION AF357220
KEYWORDS AF357220.1 GI:13448828
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 292)
Mckie, K., Addington, T., Nguyen, T.S., Glendenning, M., Kutlar, F. and
Kutlar, A.
Detection of TATA box TA repeat region [6(TA)repeat] of human
bilirubin UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in an
African American individual
Unpublished
JOURNAL 2 (bases 1 to 292)
REFERENCE 2 (bases 1 to 292)
Mckie, K., Addington, T., Nguyen, T.S., Glendenning, M., Kutlar, F. and
Kutlar, A.
Submitted (07-MAR-2001) Medicine/Hemoglobin DNA Laboratory, Medical
College of Georgia, 15th Street, Augusta, GA 30912, USA
location/Qualifiers
1..292
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2"
/map="2q37"
/cell_type="WBC"
/tissue_type="whole blood"
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/note="most common type of repeat; 6(TA)"
28..42
/gene="UGT1"
/note="polymorphic region"
38..39
/rpt_type=tandem
/rpt_unlc=ta
65..>292
/gene="UGT1"
/product="bilirubin UDP-glucuronosyltransferase 1"
81..>292
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/codon_start=1
/product="bilirubin UDP-glucuronosyltransferase 1"
/protein_id="AAK27223.1"
/db_xref="GI:13448829"
/translation="MAVESOGGPRVLLGGLCVLPVYVSHAKILLIPVDSHWLML
GAIQQLQGSGHEIVVLAIPDASLYIRNG"

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BASE COUNT 60 a 69 c 95 g 68 t
ORIGIN

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Query Match 100.0%; Score 21; DB 89; Length 292;
Best Local Similarity 100.0%; Pred. No. 0.72;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 ccaactggatcaacagatct 21
DB 186 ccaactggatcaacagatct 166

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RESULT 3
AF352795/c
LOCUS 531 bp DNA PRI 10-APR-2001
DEFINITION Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1A1) gene,
UGT1A1*1 allele, partial cds.
ACCESSION AF352795
VERSION AF352795.1 GI:13569708
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS McKie,K., Kutlar,F., Glendenning,M. and Kutlar,A.
TITLE 7(TTA) repeat polymorphism of the TATA box of human bilirubin
UDP-glucuronosyltransferase 1-1(UGT1A1*) gene in a patient with
sickle cell anemia + high bilirubinemia
JOURNAL Unpublished
2 (bases 1 to 531)
REFERENCE McKie,K., Kutlar,F., Glendenning,M. and Kutlar,A.
AUTHORS Direct Submission
TITLE Submitted (23-FEB-2001) Medicine/Hemoglobin DNA Laboratory, Medical
College of Georgia, 15th St., AC-1000, Augusta, GA 30912, USA
FEATURES
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location/Qualifiers
1..531
/organism="Homo sapiens"
/db_xref="taxon:9606"
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/map="2q37"
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/tissue_type="blood"
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1..313
/gene="UGT1A1"
279..295
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279..292
/note="repeat polymorphism compared to UGT1A1 sequence
presented in Genbank Accession Number AF180372; contains 7
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bilirubinemia"
/rpt_type=tandem
/rpt_unit=ta
314..531
/gene="UGT1A1"
/product="bilirubin UDP-glucuronosyltransferase 1-1"
334..531
/gene="UGT1A1"
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/product="bilirubin UDP-glucuronosyltransferase 1-1"
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/db_xref="GI:13569709"
/translation="MAVESGGGRPLVGLLGVLPVVSRRKILLIPVDSHWSML
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BASE COUNT 120 a 121 c 137 g 153 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 0.73;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 439 CCACGGGATCAACAGTACT 419

DEFINITION Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1) gene,
UGT1*1 allele, partial cds.
ACCESSION AF180372
VERSION AF180372.1 GI:6010649
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Kutlar,F., Stromek,E., Leitner,C., Nechtman,J. and Kutlar,A.
TITLE Detection of the TATA box polymorphism of the human bilirubin
UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in a patient with
sickle cell anemia
JOURNAL Unpublished
2 (bases 1 to 541)
REFERENCE Kutlar,F., Stromek,E., Leitner,C., Nechtman,J. and Kutlar,A.
AUTHORS Direct Submission
TITLE Submitted (24-JUG-1999) Medicine, Hematology/Oncology-Sickle Cell
Center, Medical College of Georgia, 15th Street, AC-1000, Augusta,
GA 30912, USA
FEATURES
source
location/Qualifiers
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/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2"
/map="2q37"
/cell_type="white blood cells"
/tissue_type="whole blood"
1..541
/gene="UGT1"
/note="GNT1; UGT1A1"
/allele="UGT1*1"
1..313
/gene="UGT1"
226
/gene="UGT1"
279..288
/replace="t"
/note="polymorphic region"
/rpt_type=tandem
/rpt_unit=ta
279..291
/gene="UGT1"
/note="Ritter,J.R., et al., 1992, J. Biol. Chem.,
267:3257-3261"
314..541
/gene="UGT1"
/product="bilirubin UDP-glucuronosyltransferase 1-1"
330..541
/gene="UGT1"
/note="UDP glucosyltransferase 1"
/codon_start=1
/product="bilirubin UDP-glucuronosyltransferase 1-1"
/protein_id="AAK01205.1"
/db_xref="GI:6010650"
/translation="MAVESGGGRPLVGLLGVLPVVSRRKILLIPVDSHWSML
GAIOQLQORGEHYIVLAPDASLYTRG"
BASE COUNT 123 a 124 c 141 g 153 t
ORIGIN

Query Match 100.0%; Score 21; DB 89; Length 541;
Best Local Similarity 100.0%; Pred. No. 0.73;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 435 CCACGGGATCAACAGTACT 415

RESULT 4
AF180372/c
LOCUS 541 bp DNA PRI 05-OCT-1999

RESULT 5
AF110194/c

LOCUS AF110194 918 bp DNA PRI 02-JAN-2001
 DEFINITION Homo sapiens chromosome 2 UDP-glucuronosyltransferase (UGT1A1)
 ACCESSION AF110194
 VERSION AF110194
 KEYWORDS AF110194.1 GI:12002134
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 918)
 AUTHORS Gullermette, C.
 TITLE Direct Submission
 JOURNAL Submitted (01-DEC-1998) Center for Cancer Research, MIT, 77 Massachusetts Avenue, E17-540, Cambridge, MA 02139, USA
 FEATURES
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 /db_xref="taxon:9606"
 /chromosome="2"
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 /gene="UGT1A1"
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 /note="22..918; 1A1 variant allele; L233R"
 /allele="UGT1A1*33"
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 /codon_start=1
 /product="UDP-glucuronosyltransferase"
 /protein_id="AA03197.1"
 /db_xref="GI:12002135"
 /translation="MAVESQGRPLVGLLCEVGPVSHAKILLIPDGSNLSML
 GAIQLOQRHEIVLAPDASLYTRDGAFTLTTPVPPQREDEVESVSLGNVFN
 DSFLQRIKTKYKKIKKDSAMLSGCSHLNKKELMASLSSFDVMTDPLPCSPY
 AQLSLPVEFLHAPCSLEFEATOCNPFSYPRPLSSSDMTFLQRYVNMILAFS
 QNFLCDVYSYATRASEFLQREYVQDLSSASVWLFPSDVKDPRPIMPNNVFG
 GINCHQNPESQVIGVG"
 BASE COUNT 196 a 245 c 235 g 242 t
 ORIGIN
 Query Match 100.0%; Score 21; DB 88; Length 918;
 Best Local Similarity 100.0%; Pred. No. 0.74;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

TITLE A novel complex locus ugt1 encodes human bilirubin, phenol, and other UDP-glucuronosyltransferase isozymes with identical carboxyl terminal
 JOURNAL J. Biol. Chem. 267 (5), 3257-3261 (1992)
 MEDLINE 92147680
 FEATURES
 source
 Location/Qualifiers
 1..1190
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /tissue-type="liver"
 /tissue-lib="cosmid"
 /lab_host="XLI-Blue"
 32..46
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 32..46
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 69..948
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 /number=1
 69..1190
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 85..>948
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 /codon_start=1
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 /protein_id="AA061248.1"
 /db_xref="GI:340132"
 /translation="MAVESQGRPLVGLLCEVGPVSHAKILLIPDGSNLSML
 GAIQLOQRHEIVLAPDASLYTRDGAFTLTTPVPPQREDEVESVSLGNVFN
 DSFLQRIKTKYKKIKKDSAMLSGCSHLNKKELMASLSSFDVMTDPLPCSPY
 AQLSLPVEFLHAPCSLEFEATOCNPFSYPRPLSSSDMTFLQRYVNMILAFS
 QNFLCDVYSYATRASEFLQREYVQDLSSASVWLFPSDVKDPRPIMPNNVFG
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 949..1190
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 /note="does not fit consensus"
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 BASE COUNT 261 a 290 c 286 g 353 t
 ORIGIN
 Query Match 100.0%; Score 21; DB 97; Length 1190;
 Best Local Similarity 100.0%; Pred. No. 0.74;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

BASE COUNT	ORIGIN	COMMENT	FEATURES	source
893 a	935 c	803 g	950 t	
<p> Direct Submission Submitted (04-SEP-1996) to the DDBJ/EMBL/GenBank databases. Hisao Ueyama, Shiga University of Medical Science, Department of Medical Biochemistry, Seta, Otsu, Shiga 520-21, Japan (Tel:077-548-2162, Fax:077-548-2164) </p> <p> Sequence updated (08-Jan-1997) by: Hisao Ueyama. </p> <p> Location/Qualifiers </p> <p> 1..3341 </p> <p> /organism="Homo sapiens" /db_xref="taxon:9606" /chromosome="2q37" </p> <p> misc_feature 2538..2544 /note="XRE" </p> <p> misc_feature 2606..2610 /note="XRE" </p> <p> misc_feature 3088..3097 /note="E-box" </p> <p> misc_feature 3101..3113 /note="HNF-1 site" </p> <p> CAAT_signal 3125..3129 TATA_signal 3149..3153 exon 3177..3341 /number=1 </p> <p> gene 3192..3341 /gene="UGT1*1" </p> <p> CDS 3192..3341 /gene="UGT1*1" /codon_start=1 /product="bilirubin UDP-glucuronosyltransferase 1" /protein_id="BAA25600.1" /db_xref="GI:3059177" /translation="NAVESQCRPIVIGLLCLVLPVYSHAKILILIPVDSGSHLSMLGATQD" </p>				

[illegible]

AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (13-AUG-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
REFERENCE 5 (bases 1 to 68770)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (22-OCT-1999) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
REFERENCE 6 (bases 1 to 68770)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (21-DEC-1999) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT On Aug 13, 1999 this sequence version replaced gi:4337256.

Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: <http://genome.wustl.edu/gsc>
 Contact: saplens@watson.wustl.edu
 Summary Statistics
 Center project name: H_NH0154124

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality > 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
 Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
 The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P.Y., Zhao, B., Frenken, B., Tatenio, M., Catanesse, J.J. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)
 VECTOR: pBACpac.6

NEIGHBORING SEQUENCE INFORMATION:
 The clone sequenced to the left is RP11-332L11, 200 bp overlap. Actual start of this clone is at base position 8614 of RP11-332L11; actual end is at base position 68770 of RP11-154L24.

The clone RP11-154L24 contains a tandem repeat from base positions 38234 to 39039, this region contains some low quality data. The assembly is consistent with the restriction digest information.

FEATURES
SOURCE
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 3. 192
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 repeat_region
 221. 528

/rpt_family="Alu"
 529. 1634
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 1638. 1878
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 repeat_region
 2079. 2271
 /rpt_family="MERL_type"
 repeat_region
 2276. 2336
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 4683. 4806
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 /rpt_family="L1"
 repeat_region
 11196. 11469
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 repeat_region
 11540. 11654
 /rpt_family="L1"
 repeat_region
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 13150. 13171
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 13338. 13497
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 14727. 14848
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 repeat_region
 15890. 16244
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 16380. 16723
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 17483. 17560
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 17655. 17747
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 19122. 19636
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 19637. 19662
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 repeat_region
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 repeat_region
 21441. 21162
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 25777. 25824
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 repeat_region
 28178. 28232
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repeat_region 31268..31662
repeat_region 32195..32390
repeat_region 33607..33760
repeat_region 33835..34127
repeat_region 34167..34283
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repeat_region 35247..35719
repeat_region 35720..35762
repeat_region 35763..35909

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Query Match 100.0% Score 21 DB 86 Length 68770:
Best Local Similarity 100.0% Pred. No. 0.81:
Matches 21: Conservative 0; Mismatches 0; Indels 0; Gaps 0:

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QY 1 cccactggatcaacagatct 21
Db 66050 CCACGTGATCAACAGATCT 66070

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RESULT 10
AC026497 176619 bp DNA HTG 22-MAR-2000
LOCUS Homo sapiens chromosome 11 clone RP11-689A10 map 11, *** SEQUENCING
DEFINITION
IN PROGRESS ***, 43 unordered pieces.
ACCESSION AC026497.1 GI:7283186
VERSION
KEYWORDS HTG: HTGS_PHASE1.
SOURCE human.
ORGANISM Homo sapiens

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REFERENCE
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abrahams, H., Allen, N.,
Anderson, S., Baldwin, J., Batra, N., Bastien, V., Bedalov, F.,
Boguski, M., Brown, A., Brown, A., Burt, G.,
Campbell, A., Castle, A., Chao, Y., Colangelo, M., Collins, S.,
Corry, A., Cooke, P., Dear, K., Delellano, K., Dewar, K., Diaz, J.S.,
Dodge, S., Domingo, M., Doyle, M., Fereira, P., Fitzhugh, W., Gage, D.,
Gale, J., Gardner, S., Ginde, S., Goette, M., Graham, L.,
Grand, P., Grant, G., Hagos, B., Heath, A., Horton, L.,
Klein, J., Lacombe, K., Lamazares, R., Landers, T., Lebeck, J.,
Levine, R., Liu, C., Liu, G., Locke, K., MacDonald, P., Margulis, A.,
McCarthy, M., McGraw, P., McGuire, A., McKernan, K., McPherson, R.,
Meldrum, J., Meneilly, K., Miron, T., Miranda, C., Minger, A., Morrow, J.,
Murphy, T., Nayler, J., Norman, C.H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Oliver, T.M., Oliver, J., Peterson, K., Pierre, N.,
Pless, C., Pollard, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,

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TITLE JOURNAL COMMENT

Strange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Tessier, S., Theodore, J., Tjelle, A., Travers, M., Triggillo, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J.,
Young, G., Zainoun, J., Zimmer, A. and Zody, M.
Direct Submission
Submitted (22-MAR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu

Project Information

Center project name: 16277
Center clone name: 689A10

* NOTE: This is a 'working draft' sequence. It currently
* consists of 43 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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1 1333: contig of 1333 bp in length
1334 1433: gap of 100 bp
1434 2787: contig of 1354 bp in length
2788 2887: gap of 100 bp
2888 3998: contig of 1111 bp in length
3999 4098: gap of 100 bp
4099 5777: contig of 1679 bp in length
5778 5877: gap of 100 bp
5878 7685: contig of 1808 bp in length
7686 7785: gap of 100 bp
7786 8176: contig of 391 bp in length
8177 8276: gap of 100 bp
8277 9926: contig of 1650 bp in length
9927 10026: gap of 100 bp
10027 11498: contig of 1473 bp in length
11499 11599: gap of 100 bp
11600 13616: contig of 2017 bp in length
13617 13716: gap of 100 bp
13717 16073: contig of 2357 bp in length
16074 16173: gap of 100 bp
16174 18510: contig of 2337 bp in length
18511 18610: gap of 100 bp
18611 20495: contig of 1885 bp in length
20496 20595: gap of 100 bp
20596 23548: contig of 2953 bp in length
23549 23648: gap of 100 bp
23649 25910: contig of 2262 bp in length
25911 26010: gap of 100 bp
26011 28060: contig of 2050 bp in length
28061 28160: gap of 100 bp
28161 30134: contig of 1974 bp in length
30135 30234: gap of 100 bp
30235 32646: contig of 2412 bp in length
32647 32746: gap of 100 bp
32747 35096: contig of 2350 bp in length
35097 35196: gap of 100 bp
35197 37966: contig of 2500 bp in length
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37967 40539: contig of 2743 bp in length
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* 65048 69431: contig of 4384 bp in length
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* 69532 74167: contig of 4636 bp in length
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* 74268 78188: contig of 3921 bp in length
* 78189 78288: gap of 100 bp
* 78289 81664: contig of 3376 bp in length
* 81665 81764: gap of 100 bp
* 81765 85169: contig of 3405 bp in length
* 85170 85269: gap of 100 bp
* 85270 90445: contig of 5176 bp in length
* 90446 90545: gap of 100 bp
* 90546 95008: contig of 4463 bp in length
* 95009 95108: gap of 100 bp
* 95109 100587: contig of 5479 bp in length
* 100588 100687: gap of 100 bp
* 100688 105884: contig of 5197 bp in length
* 105885 105984: gap of 100 bp
* 105985 111486: contig of 5502 bp in length
* 111487 111586: gap of 100 bp
* 111587 117781: contig of 6195 bp in length
* 117782 117881: gap of 100 bp
* 117882 125343: contig of 7462 bp in length
* 125344 125443: gap of 100 bp
* 125444 132508: contig of 7065 bp in length
* 132509 132608: gap of 100 bp
* 132609 141190: contig of 8582 bp in length
* 141191 141290: gap of 100 bp
* 141291 152556: contig of 11266 bp in length
* 152557 152656: gap of 100 bp
* 152657 163590: contig of 10934 bp in length
* 163591 163690: gap of 100 bp
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/db_xref="taxon:9606"
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2888..3998
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Best Local Similarity 100.0% Pred. No. 0.83; 0; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0;

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DB 89296 CCACGTGGATCACAGTATCT 89316
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RESULT 11
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LOCUS AF297093/c
DEFINITION Homo sapiens UGT1 gene locus, complete sequence.
ACCESSION AF297093
VERSION AF297093.1 GI:11118740
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE
1 (bases 1 to 198872)
Owens,I.S., Gong,Q., Cho,J.W., Huang,T., Potter,C., Gholami,N.,
Basu,N.K., Kubota,S., Carvalho,S. and Pennington,M.W.
Thirteen UDP glucuronosyltransferase genes encoded at the human
UGT1 locus

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JOURNAL Unpublished
2 (bases 1 to 198872)
REFERENCE
Owens,I.S., Gong,Q., Cho,J.W., Huang,T., Potter,C., Gholami,N.,
Basu,N.K., Kubota,S., Carvalho,S. and Pennington,M.W.
Direct Submission
Submitted (10-AUG-2000) Heritable Disorders Branch, NIH-NICHD, 9000
Rockville Pike, Bethesda, MD 20892, USA

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FEATURES

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 LCHRFVFNALIEISLILQTPYTHYDLSHLSHLLRDPVLDYPRVYVNMFTIGIN
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 EYLGPSVLYRGFPCSLFETRSPPDVSTIPRCYKSDHMFQSNVAFNLVNLLE
 PYLFYCLFSKYEELASAVLKRDVDITLTKOVSWILARDEVLLEYPRVNMVNLGG
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Best Local Similarity 100.0%; Pred. No. 0.83; Length 198872;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ccactggatcacagatct 21
DB 175148 CCACGTGGATCAACAGATCT 175128

RESULT 12
AY029169 1602 bp mRNA PRI 16-APR-2001
LOCUS Macaca mulatta UDP-glucuronosyltransferase UGT1A01 mRNA, complete cds
ACCESSION AY029169
VERSION AY029169.1 GI:13641264
KEYWORDS
SOURCE
ORGANISM
Thesius monkey.
Macaca mulatta
Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea; Cercopithecinae; Macaca.
REFERENCE
AUTHORS Dean, B.J., Zhao, S. and King, C.
TITLE UDP-glucuronosyltransferase 1A isoform Cloned from Male Rhesus Monkey Liver
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1602)
AUTHORS Dean, B.J., Zhao, S. and King, C.
TITLE Direct Submission
JOURNAL Submitted (02-APR-2001) Drug Metabolism, Merck & Co. Inc., Mail Drop RY80F-109, PO Box 2000, Rahway, NJ 07065, USA

FEATURES
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LFGQMDNAKRMETKAGVTLVLEMTSEDEENLAKAVINDSKYENIMHLSLHKOR
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BASE COUNT 383 a 417 c 390 g 412 t
ORIGIN

Query Match
Best Local Similarity 92.4%; Score 19.4; DB 91; Length 1603;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ccactggatcacagatct 21
DB 106 CCACGTGGATCAACAGATCT 86

RESULT 13
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LOCUS Macaca fascicularis UDP-glucuronosyltransferase UGT1A01 mRNA, complete cds.
ACCESSION AF104339
VERSION AF104339.1 GI:6537143
KEYWORDS
SOURCE
ORGANISM
craab-eating macaque.
Macaca fascicularis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea; Cercopithecinae; Macaca.
REFERENCE
AUTHORS Albert, C., Vallee, M., Beaudry, C., Belanger, A. and Hum, D.W.
TITLE 1 (bases 1 to 1647)
JOURNAL Submitted (05-NOV-1998) Molecular Endocrinology, Laval University, 2705 Laurier Boulevard, Sainte-Foy, Que G1V4G2, Canada

FEATURES
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BASE COUNT 393 a 431 c 409 g 414 t
ORIGIN

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Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ccactggatcacagatct 21
DB 143 CCACGTGGATCAACAGATCT 123

RESULT 14
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LOCUS Homo sapiens chromosome 16p24.3 clone RP4-754F23, WORKING DRAFT
DEFINITION
SEQUENCE 35 ordered pieces.
ACCESSION AC007905
VERSION AC007905.2 GI:10280722
KEYWORDS
SOURCE HTG; PHASE2; HTGS; DRAFT.
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 104480)
AUTHORS Krenmadiotis, G., Gardner, A.E., Callen, D.F. and Sutherland, G.R.

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

Large Scale Sequencing of the Chromosome 16 region q24.3
(bases 1 to 104480)
Kremmidiotis, G., Gardner, A.E., Callen, D.F. and Sutherland, G.R.
Submitted (24-JUN-1999) CytoGenetics & Molecular Genetics, Women's
& Children's Hospital, 72 King William Rd, Adelaide, SA 5006,
Australia

COMMENT

On Sep 23, 2000 this sequence version replaced g1:5174819.

Genome Centre : CytoGenetics & Molecular Genetics
Centre Code : CMGCH

Website: <http://www.wch.sa.gov.au/labmedic/genetics/sequencing.html>

* NOTE: This is a 'working draft' sequence. It currently
* consists of 35 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.

* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

1 1479: contig of 1479 bp in length
1480 1487: gap of unknown length
1488 6699: contig of 5212 bp in length
6700 6707: gap of unknown length
6708 10905: contig of 4198 bp in length
10906 10912: gap of unknown length
10913 11733: contig of 821 bp in length
11734 11740: gap of unknown length
11741 12602: contig of 862 bp in length
12603 12609: gap of unknown length
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13857 13863: gap of unknown length
13864 20018: contig of 6155 bp in length
20019 20025: gap of unknown length
20026 21942: contig of 1917 bp in length
21943 21949: gap of unknown length
21950 23920: contig of 1971 bp in length
23921 23927: gap of unknown length
23928 24860: contig of 933 bp in length
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29293 29779: gap of unknown length
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32412 32418: gap of unknown length
32419 33145: contig of 727 bp in length
33146 33152: gap of unknown length
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42182 45604: contig of 3423 bp in length
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82464 82469: gap of unknown length
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84167 84173: gap of unknown length
84174 84822: contig of 649 bp in length
84823 84829: gap of unknown length
84830 99511: contig of 14682 bp in length
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101088 101094: gap of unknown length
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103317 103323: gap of unknown length
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FEATURES

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/db_xref="taxon:9606"
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ORIGIN

Query Match

Best Local Similarity 84.8%; Score 17.8; DB 60; Length 104480;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 cccctggagtcacagctatct 21
|||||

DB 20902 CCACCTGGATCAACATATTT 20922

RESULT 15

HSDD137K2/c

DEFINITION

Human DNA sequence from clone RPI-137K2 on chromosome 6q25.1-25.3.
(contains KIAA1235), ESTs, STSS, GSSs and two putative Cpg Islands,
complete sequence.

ACCESSION

AL049820

VERSION

AL049820.23 GI:8247261

KEYWORDS

HTG; B120; CPG Island; KIAA1235.

SOURCE

human.

ORGANISM

Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE

1 (bases 1 to 187507)

AUTHORS

Sycamore, N.

TITLE

Direct Submission

JOURNAL

Submitted (18-JUL-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humanyes@sanger.ac.uk
requests: clonerequests@sanger.ac.uk

COMMENT

On Jun 4, 2000 this sequence version replaced g1:8018160.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.

This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated human repeat sequence elements (e.g. Alu). Where the
sequence is ambiguous, there is an annotation using the 'unsure'
feature key.

The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:

EMBL, SWI, SWISSPROT, Tr, TREMBL, Wp, WORMPEP, Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/RGP/chr6> RPI-137K2 is from the library RPCI-1 constructed at the Roswell Park Cancer Institute by the group of Pletzer de Jong. For further details see <http://Dacpac.med.buffalo.edu/> VECTOR: pCYPAC2

This sequence is the entire insert of clone RPI-137K2 The true right end of clone RPI-80E10 is at 2741 in this sequence.

FEATURES

source

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complement(3848..4325)
/note="match: GSS: Em:B71907"
5108. .5417
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6157. .6278
/note="MIR repeat: matches 15. .153 of consensus"
6589. .6661
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6663. .6727
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6768. .7057
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7142. .7944
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7589. .7889
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9589. .9699
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9826. .10165
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10580. .10752
/note="AluY repeat: matches 132. .302 of consensus"
10795. .10905
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10975. .11081
/note="AluY repeat: matches 14. .125 of consensus"
11763. .11796
/note="L17 copies 2 mer tt 82% conserved"
11797. .12023
/note="AluSg/X repeat: matches 86. .312 of consensus"
12067. .12348
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14350. .14877
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14357. .14684
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14623. .14676
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14677. .14870
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15034. .15291
/note="LIMB3 repeat: matches 5921. .6181 of consensus"
16601. .16797
/note="MIR repeat: matches 14. .219 of consensus"
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repeat_region 17517. 17556
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repeat_region 18805. 18879
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repeat_region 19765. 20073
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23350. .23846
/note="MER6 repeat: matches 1. .865 of consensus"
repeat_region 23847. 24019
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24447. 24872
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24438. 24762
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24739. 25035
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25108. 25407
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25938. 26203
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26263. 26352
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29780. 30137
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29812. 30102
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33258. 33283
/note="LIMB1 repeat: matches 5549. .6053 of consensus"
33823. 33107
/note="AluSg repeat: matches 1. .297 of consensus"
33581. 33709
/note="FLAM repeat: matches 1. .128 of consensus"
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34949. 34972
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36527. 36791
/note="L2 repeat: matches 2159. .2428 of consensus"
36931. 37354
/note="MIR repeat: matches 57. .504 of consensus"
38691. 38991
/note="AluX repeat: matches 1. .296 of consensus"

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repeat_region 46214..46442
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repeat_region 46533..46827
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repeat_region 46550..46747
/note="3 copies 66 mer 74% conserved"
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/note="6 copies 42 mer 66% conserved"
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/note="19 copies 13 mer 61% conserved"
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Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 ccactggatcaacagttatct 21
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DB 118886 CCACGTGCATCAGCAGTATCT 118866

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Search completed: July 25, 2001, 05:16:58
Job time: 9221 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 02:43:17 ; Search time 1290.33 Seconds

(without alignments)
251736 Million cell updates/sec

Title: US-09-142-095-1

Perfect score: 21

Sequence: 1 aaagtaactccctgctacactt 21

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1344157 seqs, 7733874588 residues

Total number of hits satisfying chosen parameters: 2688314

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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6: gb_in3:*
7: gb_cm:*
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13: gb_pi2:*
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17: em_da2:*
18: em_fun:*
19: em_hrgo_hum:*
20: em_hrgo_inv:*
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22: em_hrg_hum1:*
23: em_hrg_hum2:*
24: em_hrg_hum3:*
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96: gb_pr12:*
97: gb_pr13:*
98: em_da3:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	100.0	21	9 A65500	A65500 Sequence 1
2	21	100.0	208	89 AF135471	AF135471 Salimiri b
3	21	100.0	234	89 AF135462	AF135462 Pan penis
4	21	100.0	234	89 AF135463	AF135463 Pan trogl
5	21	100.0	531	89 AF352795	AF352795 Homo sapi
6	21	100.0	541	89 AF180372	AF180372 Homo sapi
7	21	100.0	620	9 A65504	A65504 sequence 5
8	21	100.0	3341	91 D87674	D87674 Homo sapien

c 9 21 100.0 56770 86 AC006985
 c 10 21 100.0 176619 70 AC026497
 c 11 21 100.0 198872 89 AF297093
 c 12 18.4 87.6 136868 86 AC008278
 c 13 17.8 84.8 7218 91 AP000293
 c 14 17.8 84.8 46831 77 AC087204
 c 15 17.8 84.8 47972 90 AL391843
 c 16 17.8 84.8 47000 88 AC065278
 c 17 17.8 84.8 70068 77 AC090246
 c 18 17.8 84.8 100000 91 AP000043
 c 19 17.8 84.8 100000 91 AP000111
 c 20 17.8 84.8 100000 91 AP000187
 c 21 17.8 84.8 105733 75 AC074136
 c 22 17.8 84.8 125428 88 AC073898
 c 23 17.8 84.8 139493 65 AC018537
 c 24 17.8 84.8 145540 64 AC015962
 c 25 17.8 84.8 145540 64 AC015962
 c 26 17.8 84.8 153234 70 AC026947
 c 27 17.8 84.8 156486 62 AC016557
 c 28 17.8 84.8 156771 82 AL590223
 c 29 17.8 84.8 165987 66 AC021201
 c 30 17.8 84.8 167195 87 AC009044
 c 31 17.8 84.8 170916 76 AC079414
 c 32 17.8 84.8 173985 76 AC084085
 c 33 17.8 84.8 174788 82 AP000756
 c 34 17.8 84.8 175393 73 AC068412
 c 35 17.8 84.8 175587 77 AC087616
 c 36 17.8 84.8 176107 83 AP002393
 c 37 17.8 84.8 176317 62 AC01468
 c 38 17.8 84.8 182815 62 AC011386
 c 39 17.8 84.8 183279 75 AC079014
 c 40 17.8 84.8 191602 73 AC068920
 c 41 17.8 84.8 340000 91 AP001716
 c 42 17.4 82.9 1073 53 C88063X
 c 43 17 81.0 29605 12 AB020747
 c 44 17 81.0 105223 12 AC007399
 c 45 17 81.0 156970 74 AC073330

ALIGNMENTS

RESULT 1
 A65500 21 bp DNA
 LOCUS Sequence 1 from Patent WO9732042.
 ACCESSION A65500
 VERSION A65500.1 GI:4531235
 KEYWORDS
 SOURCE unidentified.
 ORGANISM unidentified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Burchell, B.
 TITLE DRUG TRIAL ASSAY SYSTEM
 JOURNAL Patent: WO 9732042-A 1 04-SEP-1997;
 UNTV DUNDEE (GB)
 COMMENT Other publication AU 2224197 19970916.
 FEATURES
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QY 1 aagtgactccctgactct 21
 DB 1 AAGTGAACCTCCCTGCTACTCT 21

RESULT 2
 AF135471 208 bp DNA PRI 21-NOV-1999
 LOCUS Salimiri boliviensis UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
 DEFINITION promoter region and partial cds.
 ACCESSION AF135471
 VERSION AF135471.1 GI:6456559
 KEYWORDS Bolivian squirrel monkey.
 SOURCE Salimiri boliviensis
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Cebinae;
 Salimiri.
 REFERENCE 1 (bases 1 to 208)
 AUTHORS Hall, D., Ybazeza, G., Destro-Biscol, G., Petzl-Erler, M.L. and Di Rienzo, A.
 TITLE Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates
 JOURNAL Pharmacogenetics (1999) In press
 AUTHORS Ybazeza, G., Hall, D. and Di Rienzo, A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th street, Chicago, IL 60637, USA
 FEATURES
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QY 1 aagtgactccctgactct 21
 DB 3 AAGTGAACCTCCCTGCTACTCT 23

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 DEFINITION promoter region and partial cds.
 ACCESSION AF135462
 VERSION AF135462.1 GI:6456541
 KEYWORDS pygmy chimpanzee.
 SOURCE Pan paniscus
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Pan.
 REFERENCE 1 (bases 1 to 234)
 AUTHORS Hall, D., Ybazeza, G., Destro-Biscol, G., Petzl-Erler, M.L. and Di Rienzo, A.
 TITLE Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates
 JOURNAL Pharmacogenetics (1999) In press

REFERENCE 2 (bases 1 to 234)
 AUTHORS Ybazaeta,G., Hall,D. and Di Rienzo,A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
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 3 aagtgacctccctgacctt 23

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 LOCUS AF135463
 DEFINITION Pan troglodytes UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
 promoter region and partial cds.
 ACCESSION AF135463
 VERSION AF135463.1 GI:6456543
 KEYWORDS chimpanzee.
 SOURCE Pan troglodytes
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Pan.
 REFERENCE 1 (bases 1 to 234)
 AUTHORS Hall,D., Ybazaeta,G., Destro-Bisol,G., Petzl-Erler,M.L. and Di
 Rienzo,A.
 TITLE Variability at the uridine diphosphate glucuronosyltransferase 1A1
 promoter in human populations and primates
 JOURNAL Pharmacogenetics (1999) In press
 AUTHORS Ybazaeta,G., Hall,D. and Di Rienzo,A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924
 E. 57th Street, Chicago, IL 60637, USA

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QY 1 aagtgacctccctgacctt 21
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RESULT 5
 AF352795 531 bp DNA PRI 10-APR-2001
 LOCUS AF352795
 DEFINITION Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1A1) gene,
 UGT1A1*1 allele, partial cds.
 ACCESSION AF352795
 VERSION AF352795.1 GI:13569708
 KEYWORDS human.
 SOURCE Homo sapiens
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 REFERENCE 1 (bases 1 to 531)
 AUTHORS McKie,K., Kutlar,F., Glendinning,M. and Kutlar,A.
 TITLE 7(TTA) repeat polymorphism of the TATA box of human bilirubin
 UDP-glucuronosyltransferase 1-1(UGT1A1*) gene in a patient with
 sickle cell anemia + high bilirubinemia
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 531)
 AUTHORS McKie,K., Kutlar,F., Glendinning,M. and Kutlar,A.
 TITLE Direct Submission
 JOURNAL Submitted (23-FEB-2001) Medicine/Hemoglobin DNA Laboratory, Medical
 College of Georgia, 15th St., NC-1000, Augusta, GA 30912, USA

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 ta repeats frequently detected in patients with high
 bilirubinemia"
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 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 DB 185 AAGTGAACCTCCTGCTACCTT 205

RESULT 6

LOCUS AF180372 541 bp DNA PRI 05-OCT-1999
 DEFINITION Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1) gene,
 UGT1*1 allele, partial cds.
 ACCESSION AF180372
 VERSION AF180372.1 GI:6010649
 KEYWORDS
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 541)
 Kuttar, F., Stromek, E., Leitner, C., Nechtman, J. and Kuttar, A.
 TITLE Detection of the TATA box polymorphism of the human bilirubin
 JOURNAL UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in a patient with
 sickle cell anemia
 2 (bases 1 to 541)
 Kuttar, F., Stromek, E., Leitner, C., Nechtman, J. and Kuttar, A.
 REFERENCE Unpublished
 AUTHORS Direct Submission
 JOURNAL Submitted (24-AUG-1999) Medicine, Hematology/Oncology-Sickle Cell
 Center, Medical College of Georgia, 15th Street, Ac-1000, Augusta,
 GA 30912, USA

FEATURES
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 1..313
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 226
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 /replaces="c"
 279..288
 /note="polymorphic region"
 /rpt_type="tandem
 /rpt_unit="ta
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 /note="Ritter, J.K., et al., 1992, J. Biol. Chem.,
 267:3257-3261"
 314..>541
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 /product="bilirubin UDP-glucuronosyltransferase 1-1"
 330..>541
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 /note="UDP glucosyltransferase 1"
 /codon_start=1
 /product="bilirubin UDP-glucuronosyltransferase 1-1"
 /protein_id="AF01205.1"
 /db_xref="GI:6010650"
 /translation="MAVESGGRPRVYGLLCVIGPVVSHACKILLIFVDSHWLSTL
 GAIIDQQRHEIVYLAIPASLIRDG"

BASE COUNT 123 a 124 c 141 g 153 t
 ORIGIN

Query Match 100.0%; Score 21; DB 89; Length 541;
 Best Local Similarity 100.0%; Pred. No. 0.81;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 aagtgactccctgctacctt 21
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 DB 185 AAGTGAACCTCCTGCTACCTT 205

RESULT 7

LOCUS A65504 620 bp DNA PAT 29-MAR-1999
 DEFINITION Sequence 5 from Patent WO9732042.
 ACCESSION A65504
 VERSION A65504.1 GI:4531239
 KEYWORDS
 SOURCE unidentified.
 ORGANISM unidentified.
 unclassified.

REFERENCE 1 (bases 1 to 620)
 Burchell, B.
 AUTHORS DRUG TRIAL ASSAY SYSTEM
 TITLE Patent: WO 9732042-A 5 04-SEP-1997;
 JOURNAL UNIV DUNDEE (GB)
 Other publication AU 2224197 19970916.

COMMENT Location/Qualifiers
 source 1..620
 /organism="unidentified"
 /db_xref="taxon:32644"

BASE COUNT 157 a 127 c 151 g 185 t
 ORIGIN

Query Match 100.0%; Score 21; DB 9; Length 620;
 Best Local Similarity 100.0%; Pred. No. 0.8;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 aagtgactccctgctacctt 21
 ||||||||||||||||||||
 DB 465 AAGTGAACCTCCTGCTACCTT 485

RESULT 8

LOCUS D87674 3341 bp DNA PRI 14-APR-2000
 DEFINITION Homo sapiens gene for bilirubin UDP-glucuronosyltransferase 1,
 promoter region and partial cds.
 ACCESSION D87674
 VERSION D87674.1 GI:3059176
 KEYWORDS bilirubin UDP-glucuronosyltransferase 1.
 SOURCE Homo sapiens DNA.
 ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (sites)
 Doida, Y.
 AUTHORS Ueyama, H., Koiwai, O., Soeda, Y., Sato, H., Satoh, Y., Ohkubo, I. and
 TITLE Analysis of the promoter of human bilirubin
 JOURNAL UDP-glucuronosyltransferase gene (UGT1*1) in relevance to Gilbert's
 syndrome
 Hepatol. Res. 9, 152-163 (1997)

REFERENCE 2 (bases 1 to 3341)
 Ueyama, H.
 AUTHORS Direct Submission
 JOURNAL Submitted (04-SEP-1996) to the DDBJ/EMBL/GenBank databases. Hisao
 Ueyama, Shiga University of Medical Science, Department of Medical
 Biochemistry, Seto, Otsu, Shiga 520-21, Japan (Tel:077-548-2162,
 Fax:077-548-2164)

COMMENT Sequence updated (08-Jan-1997) by: Hisao Ueyama.
 FEATURES Location/Qualifiers
 source 1..3341

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2q37"
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2538..2544
/note="XRE"
misc_feature
2606..2610
/note="XRE"
misc_feature
3088..3097
/note="E-box"
misc_feature
3101..3113
/note="HNF-1 site"
CAAT_signal
3125..3129
TATA_signal
3149..3153
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3177..3341
/number=1
gene
3192..3341
/gene="UGT1*1"
3192..3341
/gene="UGT1*1"
/codon_start=1
/product="Bilirubin UDP-glucuronosyltransferase 1"
/db_xref="GI:3059177"
/translation="MAVESGGGRFVGLLGLVGLPVVSHAGKILLIPYDGSHWLML
GAIQOL"
BASE COUNT      893 a      695 c      803 g      950 t
ORIGIN
Query Match      100.0%; Score 21; DB 91; Length 3341;
Best local similarity 100.0%; Prid. No. 0.73;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 aagtaactccctgctacctt 21
|||||
Db      3045 AAGTGAAGTCCCTGCTACCTT 3065

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RESULT 9
AC006985/c
LOCUS AC006985 68770 bp DNA PRI 21-DEC-1999
DEFINITION Homo sapiens BAC clone RP11-154L24 from 2, complete sequence.
ACCESSION AC006985
VERSION AC006985.2 GI:5732165
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 68770)
Sulston, V.E. and Waterston, R.
Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
REFERENCE 2 (bases 1 to 68770)
AUTHORS Gattling, S., Stoneking, T. and Davidson, T.
TITLE The sequence of Homo sapiens BAC clone RP11-154L24
JOURNAL Unpublished
3 (bases 1 to 68770)
REFERENCE 3 (bases 1 to 68770)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (05-MAR-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 68770)
REFERENCE 4 (bases 1 to 68770)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (13-DEC-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 68770)
REFERENCE 5 (bases 1 to 68770)
AUTHORS Waterston, R.
TITLE Direct Submission

JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Submitted (22-OCT-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
6 (bases 1 to 68770)
Waterston, R.
Direct Submission
Submitted (21-DEC-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Aug 13, 1999 this sequence version replaced gi:14337256.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@wustl.edu
----- Summary Statistics
Center project name: H.NH0154124

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D.
Matheron, Department of Genetics, Washington University, St. Louis
MO. For additional information about the map position of this
sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RP11-11 human BAC library was made from the blood of one male
donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E.,
Tateno, M., Caranese, J.J. and de Jong, P.J. (1998) An improved
approach for construction of bacterial artificial chromosome
libraries. Genomics 51:1-8. The clone may be obtained either from
Research Genetics, Inc. (<http://www.resgen.com>) or Pletier de Jong
and coworkers at the Roswell Park Cancer Institute
(<http://bacpac.med.buffalo.edu>)
VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-332L11, 200 bp overlap.
Actual start of this clone is at base position 68134 of
RP11-332L11; actual end is at base position 68770 of RP11-154L24.

The clone RP11-154L24 contains a tandem repeat from base positions
3834 to 39039, this region contains some low quality data. The
assembly is consistent with the restriction digest information.

FEATURES
source
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/db_xref="taxon:9606"
/chromosome="2"
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/rpt_family="L1"
3..192
/rpt_family="L1"
221..528
/rpt_family="Alu"
529..1634
/rpt_family="L1"
1658..1878
/rpt_family="L1"
2079..2271
/rpt_family="MER1-type"
repeat_region
2276..2336

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4683..4806
/rpt_family="MIR"
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5131..5389
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8738..8871
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9092..9371
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repeat_region      /rpt_family="MALR"
9967..10582
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11196..11469
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13150..13171
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repeat_region      /rpt_family="L1"
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repeat_region      /rpt_family="A-rich"
19637..19662
/rpt_family="A-rich"
repeat_region      /rpt_family="A-rich"
19825..20014
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repeat_region      /rpt_family="L1"
20034..20098
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repeat_region      /rpt_family="L1"
20254..21067
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repeat_region      /rpt_family="A-rich"
21411..21182
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25258..25447
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repeat_region      30867..31015
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/rpt_family="MALR"
repeat_region      32195..32390
/rpt_family="MIR"
repeat_region      33607..33760
/rpt_family="L2"
repeat_region      33835..34127
/rpt_family="Alu"
repeat_region      34167..34283
/rpt_family="GA-rich"
repeat_region      34806..34976
/rpt_family="L2"
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repeat_region      35720..35762
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repeat_region      35763..35909

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Query Match 100.0%; Score 21; DB 86; Length 68770;
 Best Local Similarity 100.0%; Pred. No. 0.62;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 aagtgaaatccctgcctactt 21
 Db 66302 AAGTGAATCCCTGCCTACTT 66282

RESULT 10
 AC026497/C
 LOCUS 176619 bp DNA HTG 22-MAR-2000
 DEFINITION Homo sapiens chromosome 11 clone RP11-689A10 map 11, *** SEQUENCING
 IN PROGRESS ***; 43 unordered pieces.
 AC026497
 VERSION AC026497.1 GI:7283186
 KEYWORDS HTG; HTGS; PHASEL.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 176619)
 AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
 Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
 Boguski,M., Bouckgeater,B., Brown,A., Burkett,G.,
 Campiano,A., Castle,A., Choquet,Y., Colangelo,M., Collins,S.,
 Collymore,A., Cooke,P., DeRubeis,K., Dewar,K., Diaz,J.S.,
 Dodge,S., Domino,M., Doyle,M., Ferreira,P., Fitzhugh,W., Gage,D.,
 Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
 Grand-Pierre,N., Grant,G., Hags,B., Heath,J., Horton,L.,
 Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karas,A.,
 Klein,R., Larocque,K., Lamazares,R., Landers,T., Lehotzky,J.,
 Levine,R., Liu,C., Liu,G., Locke,K., Macdonald,P., Margulis,N.,
 McCarthy,M., McEwan,P., McKernan,K., McPherson,R.,
 McPhy,T., Meneses,L., Mihova,T., Miranda,C., Minger,V., Morrow,J.,
 Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
 O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
 Pleasant,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
 Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
 Stange-Rhoman,N., Stojanovic,N., Subramanian,A., Talamas,J.,
 Teste,S., Theodore,J., Tirrell,A., Travers,M., Triggillo,J.,
 Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
 Young,G., Zainoun,J., Zilmer,A. and Zody,M.

TITLE
 JOURNAL
 COMMENT
 Submitted (22-MAR-2000) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIR

Web site: <http://www.seq.wi.mit.edu>

Contact: sequence.submissions@genome.wi.mit.edu

Center project name: 689_A.10

Center clone name: 689_A.10

NOTE: This is a 'working draft' sequence. It currently consists of 43 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1333: contig of 1333 bp in length
 1334 1433: gap of 100 bp
 1434 2787: contig of 1353 bp in length
 2788 2887: gap of 100 bp
 2888 3998: contig of 1111 bp in length
 3999 4098: gap of 100 bp
 4099 5777: contig of 1678 bp in length
 5778 5877: gap of 100 bp
 5878 7685: contig of 1808 bp in length
 7686 7785: gap of 100 bp
 7786 8176: contig of 391 bp in length
 8177 8276: gap of 100 bp
 8277 9926: contig of 1650 bp in length
 9927 10026: gap of 100 bp
 10027 11499: contig of 1473 bp in length
 11500 11599: gap of 100 bp
 11600 13616: contig of 2017 bp in length
 13617 13716: gap of 100 bp
 13717 16073: contig of 2357 bp in length
 16074 16173: gap of 100 bp
 16174 18510: contig of 2337 bp in length
 18511 18610: gap of 100 bp
 18611 20495: contig of 1885 bp in length
 20496 20595: gap of 100 bp
 20596 23548: contig of 2953 bp in length
 23549 23648: gap of 100 bp
 23649 25810: contig of 2262 bp in length
 25811 26010: gap of 100 bp
 26011 28060: contig of 2050 bp in length
 28061 28160: gap of 100 bp
 28161 30134: contig of 1974 bp in length
 30135 30234: gap of 100 bp
 30235 32646: contig of 2412 bp in length
 32647 32746: gap of 100 bp
 32747 35056: contig of 2350 bp in length
 35057 35196: gap of 100 bp
 35197 37696: contig of 2500 bp in length
 37697 37796: gap of 100 bp
 37797 40538: contig of 2743 bp in length
 40539 40639: gap of 100 bp
 40640 45518: contig of 4879 bp in length
 45519 45618: gap of 100 bp
 45619 48728: contig of 3110 bp in length
 48729 48828: gap of 100 bp
 48829 52099: contig of 3271 bp in length
 52100 52199: gap of 100 bp
 52200 56747: contig of 4548 bp in length
 56748 56847: gap of 100 bp
 56848 60467: contig of 3650 bp in length
 60468 60567: gap of 100 bp
 60568 64947: contig of 4360 bp in length
 64948 65047: gap of 100 bp
 65048 69431: contig of 4384 bp in length
 69432 69531: gap of 100 bp

FEATURES

SOURCE

69532 74167: contig of 4636 bp in length
 74168 74267: gap of 100 bp
 74268 76188: contig of 3921 bp in length
 76189 78288: gap of 100 bp
 78289 81664: contig of 3376 bp in length
 81665 81764: gap of 100 bp
 81765 85169: contig of 3405 bp in length
 85170 85269: gap of 100 bp
 85270 90445: contig of 5176 bp in length
 90446 90545: gap of 100 bp
 90546 95008: contig of 4463 bp in length
 95009 95108: gap of 100 bp
 95109 100587: contig of 5479 bp in length
 100588 100687: gap of 100 bp
 100688 105884: contig of 5197 bp in length
 105885 105984: gap of 100 bp
 105985 111486: contig of 5502 bp in length
 111487 111586: gap of 100 bp
 111587 117781: contig of 6195 bp in length
 117782 117881: gap of 100 bp
 117882 125343: contig of 7462 bp in length
 125344 125443: gap of 100 bp
 125444 132508: contig of 7065 bp in length
 132509 132608: gap of 100 bp
 132609 141190: contig of 8582 bp in length
 141191 141290: gap of 100 bp
 141291 152556: contig of 11266 bp in length
 152557 152656: gap of 100 bp
 152657 163590: contig of 10934 bp in length
 163591 163690: gap of 100 bp
 163691 176619: contig of 12929 bp in length.

Location/Qualifiers

1..176619

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="11"

/map="11"

/clone="RP11-689A10"

/clone_id="RPC1-11 Human Male BAC"

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/note="assembly-fragment"

1434..2787

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2888..3998

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4099..5777

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5878..7685

/note="assembly-fragment"

7786..8176

/note="assembly-fragment"

clone_end:SP6

vector_side:right"

8277..9926

/note="assembly-fragment"

10027..11499

/note="assembly-fragment"

11600..13616

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13717..16073

/note="assembly-fragment"

16174..18510

/note="assembly-fragment"

18511..20495

/note="assembly-fragment"

20586..23548

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23649..25810

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26011..28060

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Query Match      100.0%: Score 21; DB 70; Length 176619;
Best Local Similarity 100.0%: Pred. No. 0.59;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 aagtaactcctgcctactt 21
      Db 89548 AAGTGAAGCTCCTGCTACTT 89528

RESULT 11
LOCUS      AF297093      198872 bp      DNA
DEFINITION Homo sapiens UGT1 gene locus, complete sequence.
ACCESSION  AF297093
VERSION     AF297093.1 GI:11118740
KEYWORDS
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 198872)
AUTHORS     Owens,I.S., Gong,Q., Cho,J.W., Huang,T., Potter,C., Gholami,N.,
            Basu,N.K., Kubota,S., Carvalho,S. and Pennington,M.W.
            Thirteen UDP glucuronosyltransferase genes encoded at the human
            UGT1 locus
JOURNAL     Unpublished
            2 (bases 1 to 198872)
REFERENCE   Owens,I.S., Gong,Q., Cho,J.W., Huang,T., Potter,C., Gholami,N.,
            Basu,N.K., Kubota,S., Carvalho,S. and Pennington,M.W.
            Direct Submission
TITLE       Submitted (10-AUG-2000) Heritable Disorders Branch, NIH-NICHD, 9000
            Rockville Pike, Bethesda, MD 20892, USA
FEATURES
            Location/Qualifiers
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            /note="UDP glucuronosyltransferase 1A12"
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            19970..20956
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            /note="UDP glucuronosyltransferase 1A11"
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            19970..19978
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            20105..20956
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            /pseudo
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            34133..34149
            /gene="UGT1A8"
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            34133..>187313
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182974..183193,187016..>187313)
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join(34264..35118,181788..181919,182603..182690,
182974..183193,187016..187313)
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FSLPSVFAFGAGICHYLEGACCPAPLSYPRILLFESDAPVFERVKNHINLEBHL
FCQYFSKNALEISAILQTPYAVDLSHSISIMLTDTVDLYDRPVPMPNITLGIN
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DOMDKAKMETKAGVTLNLTMTSEDLNKLAVINDKSYENIMLSLHDPREVE
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52972..52988
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52972..>187313
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182974..183193,187016..>187313)
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FSLPSVFAFGAGICHYLEGACCPAPLSYPRILLFESDAPVFERVKNHINLEBHL
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/note="UDP glucuronosyltransferase 1A13"
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/gene="UGT1A13p"
/codon_start=1
/pseudo
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/gene="UGT1A9"
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FCQYFSKNALEISAILQTPYAVDLSHSISIMLTDTVDLYDRPVPMPNITLGIN
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/gene="UGT1A7"
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/product="UDP glucuronosyltransferase 1A7"
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RSAPSLTSSNGSIDLFPSCNCRSTPKDKLYEIKESCDAVFLDPACGLVAKY
FSLPSVYPMRGTFCYHLESGNCRAPLSVPRLLGSPDAVPEKRYRNHILHEHL
FCYFFKNVLEIASILLDTPTATDLSTSTIMLRTPTVLEPKPMNMLPTGGIN
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DQDNKAKMETKAGAVTLNLEMTSEDLNALKAVINDSKYKEMINLSLSKDRPVE
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EYGLSVYLRGPRCSLEHFSRSDPVSYPDYTESDSHMFESORVAPNPLVLE
PIYPTCLRSKIEELASATLRDNDITITLOKYSVLLAYDVLVLEPRVPMNPIGIG
INCKRRKDLDSQFEAYINASGEHIVSLSGMSSELPKKAALADALGKIPOTVIM
RYTGRPSNLANNITLVKMLPONDLLGHPMRAITAGSHGVYESICNGVPMWPL
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129525..>187313
/gene="UGT1A5"
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/join(<129600..130466,181788..181919,182603..182690,
182974..183193,187016..>187313)
/gene="UGT1A5"
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182974..183193,187016..187313)
/gene="UGT1A5"
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/product="UDP glucuronosyltransferase 1A5"
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/db_xref="GI:1118746"
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Query Match      100.0%; Score 21: DB 89; Length 198872;
Best Local Similarity 100.0%; Pred. No. 0.59
Matches 21: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 1 aagtgaaactccctgactct 21
|||||
Db 174896 AAGTGAACCTCCGCTACCTT 174916

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RESULT 12

AC008278/c

AC008278 136868 bp DNA PRI 07-OCT-2000

DEFINITION Homo sapiens BAC clone RP11-422A6 from 2, complete sequence.

AC008278

AC008278.3 GI:7408138

HNC

human.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

Toward a complete human genome sequence

Sulston, J.F. and Waterston, R.

Genome Res. 8 (11), 1097-1108 (1998)

2 (bases 1 to 136868)

Du, F., Stoneking, T. and Doeber, A.

The sequence of Homo sapiens BAC clone RP11-422A6

unpublished

3 (bases 1 to 136868)

Waterston, R.H.

Direct Submission

Submitted (31-JUL-1999) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

4 (bases 1 to 136868)

Waterston, R.H.

Direct Submission

Submitted (07-APR-2000) Genome Sequencing Center, Washington

University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

On Apr 4, 2000 this sequence version replaced gi.5870295.

Genome Center

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: http://genome.wustl.edu/gsc

Contact: saplens@wustl.edu

Summary Statistics

Center project name: R_NH0422A06

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap

* *
8853 85321968 contigs of 658 bp in length
6853 7610: 7610: 100 bp
7611 7710: gap of

```

* 7711 8355: contig of 645 bp in length
* 8356 8455: gap of 100 bp
* 8456 9129: contig of 674 bp in length
* 9130 9223: gap of 100 bp
* 9223 9881: contig of 662 bp in length
* 9882 9991: gap of 100 bp
* 9992 10665: contig of 674 bp in length
* 10666 10765: gap of 100 bp
* 10766 11423: contig of 658 bp in length
* 11424 11523: gap of 100 bp
* 11524 12180: contig of 667 bp in length
* 12181 12280: gap of 100 bp
* 12281 12945: contig of 665 bp in length
* 12946 13045: gap of 100 bp
* 13046 13692: contig of 647 bp in length
* 13693 13792: gap of 100 bp
* 13793 14442: contig of 650 bp in length
* 14443 14542: gap of 100 bp
* 14543 15203: contig of 661 bp in length
* 15204 15303: gap of 100 bp
* 15304 15968: contig of 665 bp in length
* 15969 16068: gap of 100 bp
* 16069 16747: contig of 679 bp in length
* 16748 16847: gap of 100 bp
* 16848 17509: contig of 662 bp in length
* 17510 17609: gap of 100 bp
* 17610 18256: contig of 647 bp in length
* 18257 18356: gap of 100 bp
* 18357 19030: contig of 674 bp in length
* 19031 19130: gap of 100 bp
* 19131 19779: contig of 649 bp in length
* 19780 19879: gap of 100 bp
* 19880 20538: contig of 659 bp in length
* 20539 20638: gap of 100 bp
* 20639 21294: contig of 656 bp in length
* 21295 21394: gap of 100 bp
* 21395 22069: contig of 675 bp in length
* 22070 22169: gap of 100 bp
* 22170 22845: contig of 676 bp in length
* 22846 22945: gap of 100 bp
* 22946 23627: contig of 682 bp in length
* 23628 23727: gap of 100 bp
* 23728 24395: contig of 668 bp in length
* 24396 24495: gap of 100 bp
* 24496 25150: contig of 655 bp in length
* 25151 25250: gap of 100 bp
* 25251 25913: contig of 663 bp in length
* 25914 26013: gap of 100 bp
* 26014 26661: contig of 648 bp in length
* 26662 26761: gap of 100 bp
* 26762 27441: contig of 680 bp in length
* 27442 27541: gap of 100 bp
* 27542 28201: contig of 660 bp in length
* 28202 28301: gap of 100 bp
* 28302 28991: contig of 680 bp in length
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* 29092 29766: contig of 675 bp in length
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* 29867 30521: contig of 655 bp in length
* 30522 30621: gap of 100 bp
* 30622 31300: contig of 679 bp in length
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* 31401 32074: contig of 674 bp in length
* 32075 32174: gap of 100 bp
* 32175 32854: contig of 680 bp in length
* 32855 32954: gap of 100 bp
* 32955 33638: contig of 685 bp in length
* 33640 33739: gap of 100 bp
* 33740 34430: contig of 691 bp in length
* 34431 34530: gap of 100 bp
* 34531 35185: contig of 655 bp in length
* 35186 35285: gap of 100 bp
* 35286 35958: contig of 673 bp in length

```

FEATURES

source

```

* 35959 36058: gap of 100 bp
* 36059 36729: contig of 671 bp in length
* 36730 36829: gap of 100 bp
* 36830 37509: contig of 680 bp in length
* 37510 37609: gap of 100 bp
* 37610 38263: contig of 654 bp in length
* 38264 38363: gap of 100 bp
* 38364 39039: contig of 676 bp in length
* 39040 39139: gap of 100 bp
* 39140 39807: contig of 668 bp in length
* 39808 39907: gap of 100 bp
* 39908 40578: contig of 671 bp in length
* 40579 40678: gap of 100 bp
* 40679 41357: contig of 679 bp in length
* 41358 41457: gap of 100 bp
* 41458 42167: contig of 710 bp in length
* 42168 42267: gap of 100 bp
* 42268 42963: contig of 696 bp in length
* 42964 43063: gap of 100 bp
* 43064 43713: contig of 650 bp in length
* 43714 43813: gap of 100 bp
* 43814 44501: contig of 688 bp in length
* 44502 44601: gap of 100 bp
* 44602 45262: contig of 661 bp in length
* 45263 45362: gap of 100 bp
* 45363 46040: contig of 678 bp in length
* 46041 46140: gap of 100 bp
* 46141 46831: contig of 691 bp in length.

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Location/Qualifiers

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/db_xref="taxon:9606"
/chromosome="8"
/map="8"

/clone="RP11-520F7"
/clone_lib="RPCT-11 Human Male BAC"

BASE COUNT 10780 a 8764 c 9356 g 11744 t 6187 others

Query Match

Best Local Similarity 84.8%; Score 17.8; DB 77; Length 46831;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 aagtgactccctcactcctt 21

Db 7326 AGCTGAGCTCCCTCCTACTT 7346

RESULT 15

AL391843/C

LOCUS

DEFINITION

sequence.

AL391843

VERSION

AL391843.13

KEYWORDS

HTG.

SOURCE

ORGANISM

human.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 47972)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AL391843 47972 bp DNA PRI 22-FEB-2001
Human DNA sequence from clone RP11-592B10 on chromosome 6, complete
sequence.
AL391843
AL391843.13 GI:13157590
HTG.
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 47972)
Sycamore, N.
Direct Submission
Submitted (22-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
Requests: clonerequests@sanger.ac.uk
On Feb 27, 2001 this sequence version replaced GI:12831877.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with

Only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality > 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; SW: SWSRPOT; Tr: TREMBL; Mp: MORNREP; information on the MORNREP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormrep. This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr6>.

Rp11-592B10 is from the library RPCI-11.3 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/dacpac/home.htm>.

VECTOR: pBAC3.6

IMPORTANT: This sequence is not the entire insert of clone Rp11-592B10. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The true left end of clone Rp11-411F9 is at 47873 in this sequence. The true right end of clone Rps-1007B16 is at 100 in this sequence.

Location/Qualifiers

FEATURES

source

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1. 47972
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/complement(156..512)
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164..366
/feature="LRR12 repeat: matches 18..192 of consensus"
332..503
/feature="match: GSS: Em:AQ542267"
427..472
/feature="LRR12 repeat: matches 189..234 of consensus"
473..549
/feature="LRR30 repeat: matches 544..620 of consensus"
480..583
/feature="MER61E repeat: matches 377..477 of consensus"
584..673
/feature="LRR12 repeat: matches 582..671 of consensus"
674..7342
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7344..7825
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7826..11300
/feature="LRR1 repeat: matches 685..4254 of consensus"
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11612..13713
/feature="LRR1 repeat: matches 4254..6304 of consensus"
13766..15206
/feature="LRR3 repeat: matches 2143..3546 of consensus"
15207..15512
/feature="AluX repeat: matches 2..308 of consensus"
15513..17628
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18106..18525

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20556..21581
/feature="MER18 repeat: matches 1..635 of consensus"
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/feature="L2 repeat: matches 2639..2695 of consensus"
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/feature="match: GSS: Em:AQ376698"
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26543..26868
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26950..27010
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27375..27869
/feature="LRR15 repeat: matches 1..483 of consensus"
27870..27942
/feature="MER90 repeat: matches 543..615 of consensus"
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/feature="match: GSS: Em:AQ82343"
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/feature="Alu repeat: matches 1..305 of consensus"
33533..33906
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35260..35321
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35346..35777
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35952..36169
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36201..36250
/feature="25 copies 2 mer ca 82% conserved"
36271..36393
/feature="FLAN_A repeat: matches 2..124 of consensus"
37540..37613
/feature="37 copies 2 mer aa 67% conserved"
38348..38398
/feature="26 copies 2 mer at 84% conserved"
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40021..40482
/feature="match: GSS: Em:AQ171081"
40102..40352
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40581..40676
/feature="48 copies 2 mer aa 64% conserved"

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/note="L1M2 repeat: matches -676. -231 of consensus"
repeat_region 42139. 42439 repeat: matches -265. -24 of consensus"
/note="L1PA15-16 repeat: matches -265. -24 of consensus"
repeat_region 42901. 43464
/note="L1PA13 repeat: matches 802. -128 of consensus"
repeat_region 43465. 43687
/note="MR30 repeat: matches 1. -230 of consensus"
repeat_region 43688. 43754
/note="L1PA13 repeat: matches 1288. -1351 of consensus"
repeat_region 43755. 46738
/note="L1PA16 repeat: matches 3178. -6153 of consensus"
repeat_region 46921. 47311
/note="L1M2 repeat: matches 1. -304 of consensus"
misc_feature complement(47070. 47394)
/note="match: GSS: Em:A053386"
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/note="match: GSS: Em:A0534016"
misc_feature 47878. 47972
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BASE COUNT 16896 a 8982 c 8503 g 13591 t
ORIGIN
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Best Local Similarity 90.58; Pred. No. 40;
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QY 1 aagtgaaactccctgctacctt 21
|||||
Db 29145 AAGTGAACCTCCTCCTT 29125
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Search completed: July 25, 2001, 05:16:42
Job time: 9205 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 04:05:07 : Search time 247.87 Seconds

(without alignments)
53.197 Million cell updates/sec

Title: US-09-142-095-1

Perfect score: 21
Sequence: 1 aagtgactccctgcactctt 21

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 730101 segs, 313950809 residues

Total number of hits satisfying chosen parameters: 1460202

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: /SIDSL/gcgdata/geneseq/geneseq/NA1980.DAT:*
2: /SIDSL/gcgdata/geneseq/geneseq/NA1981.DAT:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	100.0	21	AAAT79541	UGT1*1 gene exon 1
2	21	100.0	21	AAAT79540	Upstream DNA seque
3	16.8	80.0	237326	AAV57903	Hereditary haemoch
4	16.2	77.1	260	AAAC25716	Human secreted pro
5	16.2	77.1	867	AAAC38612	Arabidopsis thalia
6	16.2	77.1	1613	AAAC45500	CDNA encoding enzy
7	16.2	77.1	32042	AAZ09252	Human CARD-4 DNA
8	16.2	77.1	32042	AAZ09252	Human CARD-4 gene
9	15.8	75.2	957	AAO97865	TRONP1 DNA, TRePO
10	15.8	75.2	957	AAO97865	Triponema pallidum
11	15.2	72.4	537	AAAG7251	Eucalyptus grandis

12	15.2	72.4	640	18	AAV04426	Human calpain smal
13	15.2	72.4	804	18	AAV04420	Human calpain smal
14	15.2	72.4	1242	21	AACT77765	Human cancer assoc
15	15.2	72.4	1741	21	AAZ98118	Human signal pepti
16	15.2	72.4	1938	21	AAAC6514	Arabidopsis thalia
17	15.2	72.4	1940	21	AAAC39039	Arabidopsis thalia
18	15.2	72.4	2024	21	AAAC81718	Human secreted pro
19	15.2	72.4	2582	20	AAV99562	Soybean lysine ket
20	15.2	72.4	4044	21	AAAC81895	A. thaliana Srp30
21	15.2	72.4	4655	18	AAV97955	Human epididymis-s
22	15.2	72.4	235033	19	AAV57926	Hereditary haemoch
23	15	71.4	1330	21	AAAC3777	Arabidopsis thalia
24	15	71.4	14063	20	AAAZ0500	Polynucleotide seq
25	14.8	70.5	307	21	AAAC88243	Human colon cancer
26	14.8	70.5	413	21	AAAC6942	Human secreted pro
27	14.8	70.5	731	16	AAAT20482	Human gene signatu
28	14.8	70.5	1398	20	AAAZ5083	Potato tuber-speci
29	14.8	70.5	1723	17	AAAT42133	Ascorbate-free-rad
30	14.8	70.5	2161	20	AAAZ30444	Tobacco anthranila
31	14.8	70.5	2526	20	AAAZ20026	Human rad17 cell c
32	14.8	70.5	2535	20	AAAZ20032	Human rad17 cell c
33	14.8	70.5	2540	20	AAAZ20032	Human rad17 cell c
34	14.8	70.5	2543	20	AAAZ20051	Human rad17 cell c
35	14.8	70.5	2647	20	AAAZ20035	Human rad17 cell c
36	14.8	70.5	2652	17	AAAT44655	Testis-associated
37	14.8	70.5	2877	17	AAAT30085	Human Raf1 kinase
38	14.8	70.5	2877	18	AAAT61894	Human Raf-1 CDNA
39	14.8	70.5	2977	19	AAAZ0439	Human c-raf oncoge
40	14.8	70.5	2977	20	AAAZ9340	Human c-raf sequen
41	14.8	70.5	2977	20	AAAZ8137	Human c-raf-1 gene
42	14.8	70.5	2977	21	AAAZ3552	Polynucleotide #1
43	14.8	70.5	2977	21	AAAZ8554	Human c-raf DNA
44	14.8	70.5	2977	22	AAAT5126	Human c-raf CDNA
45	14.8	70.5	5907	21	AACT6910	Human OREX ORF2465

ALIGNMENTS

RESULT	ID	Score	Description
1	AAAT79541	21	UGT1*1 gene exon 1 upstream PCR primer A.
2	AAAT79541	21	Standard: DNA; 21 BP.
3	AAAT79541	21	23-JAN-1998 (first entry)
4	AAAT79541	21	UGT1*1 gene exon 1 upstream PCR primer A.
5	AAAT79541	21	Uridine diphosphate glucuronosyltransferase gene; UGT1
6	AAAT79541	21	Gilbert's syndrome; GS; unconjugated hyperbilirubinaemia;
7	AAAT79541	21	Bilirubin glucuronidation; Crigler-Najjar type 2; drug metabolism;
8	AAAT79541	21	Drug trial efficiency; screening; PCR primer; ss.
9	AAAT79541	21	Synthetic.
10	AAAT79541	21	Homo sapiens.
11	AAAT79541	21	WO9732042-A2.
12	AAAT79541	21	04-SEP-1997.
13	AAAT79541	21	03-MAR-1997; 97NO-GB00577.
14	AAAT79541	21	16-MAR-1996; 96GB-0005598.
15	AAAT79541	21	01-MAR-1996; 96GB-0004480.
16	AAAT79541	21	(UYDU-) UNIV DUNDEE.
17	AAAT79541	21	Burchell B.
18	AAAT79541	21	WPI: 1997-448702/41.
19	AAAT79541	21	Improving drug trial efficiency comprises identifying participants

PT with Gilbert's syndrome - useful as their altered drug metabolism
 PT may hinder result interpretation
 PS Claim 14; Page 12; 31pp; English.
 CC This PCR primer (with primers AAT79542-44) flanks the TATA box sequence
 CC upstream of the uridine diphosphate glucuronosyltransferase (UGT) gene
 CC 1*1 exon 1 (see AAT79540), and was used to amplify fragments of 253-255
 CC bp. This gene is known to be associated with Gilbert's syndrome (GS). GS
 CC is a mild, common form of unconjugated hyperbilirubinaemia associated
 CC with reduced bilirubin glucuronidation capacity. Analysis of the genetic
 CC basis of GS has allowed 2 forms to be identified. One is a mild form
 CC associated with a homozygous 2 bp insertion in the TATA sequence
 CC upstream of the UGT*1 exon 1, and the other is a more severe form
 CC associated with heterozygosity for a mutation which, when homozygous,
 CC causes Crigler-Najjar type 2 disease. The first form is autosomal
 CC recessive and the second is inherited dominantly. Patients suffering from
 CC GS, which is benign, may have altered metabolism of some drugs, making it
 CC difficult to determine if an effect is due to the drug or the syndrome.
 CC Drug trial efficiency would be improved if potential participants can be
 CC screened for the genetic basis of GS, and eliminated or included on
 CC basis of them possessing or not possessing GS.
 SQ Sequence 21 BP; 5 A; 7 C; 3 G; 6 T; 0 other:
 Query Match 100.0%; Score 21; DB 18; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.14; 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0;
 OY 1 aagtaacctcctgctacctt 21
 DB 1 aagtaacctcctgctacctt 21
 RESULT 2
 AAT79540
 ID AAT79540 standard; DNA; 620 BP.
 XX
 AC AAT79540:
 XX
 DT 23-JAN-1998 (first entry)
 XX
 DE Upstream DNA sequence of UGT*1 gene exon 1.
 XX
 KW uridine diphosphate glucuronosyltransferase gene; UGT;
 KW Gilbert's syndrome; GS; unconjugated hyperbilirubinaemia;
 KW bilirubin glucuronidation; Crigler-Najjar; type 2; drug metabolism;
 KW Drug trial efficiency; screening; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT protein_bind 2..8
 FT /tag- a
 FT /bound_moely- AP1
 FT GC_signal 93..102
 FT /tag- b
 FT /note- "SPL binding site"
 FT misc_feature 180..191
 FT /tag- c
 FT /note- "feature indicated in patent, but no further
 FT explanation is given"
 FT protein_bind 316..323
 FT /tag- d
 FT /bound_moely- AP3
 FT protein_bind 317..324
 FT /tag- e
 FT /bound_moely- CLBP
 FT misc_feature 366..376
 FT /tag- f
 FT /note- "feature indicated in patent, but no further
 FT explanation is given"

FT protein_bind 386..392
 FT /tag- g
 FT /bound_moely- AP1
 FT protein_bind 513..519
 FT /tag- h
 FT /bound_moely- AP1
 FT misc_feature 513..520
 FT /tag- i
 FT /note- "feature indicated in patent, but no further
 FT explanation is given"
 FT protein_bind 520..532
 FT /tag- j
 FT /bound_moely- HNF1
 FT TATA_signal 558..572
 FT /tag- k
 FT /note- "corresponds to positions -53 to -39 in patent"
 FT misc_feature 595
 FT /tag- l
 FT /note- "feature indicated in patent, but no further
 FT explanation is given"
 FT CDS 612
 FT /tag- m
 FT /codon_start- 612
 FT
 FT WO9732042-A2.
 FT
 FT PD 04-SEP-1997.
 FT
 FT XX 03-MAR-1997; 97WO-GB00577.
 FT
 FT XX 16-MAR-1996; 96GB-0005598.
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 FT PR 01-MAR-1996; 96GB-0004480.
 FT
 FT XX (UYDU-) UNIV DUNDEE.
 FT
 FT PA Burchell B;
 FT
 FT PI WPI; 1997-448702/41.
 FT
 FT DR
 FT XX
 FT PT Improving drug trial efficiency comprises identifying participants
 FT with Gilbert's syndrome - useful as their altered drug metabolism
 FT may hinder result interpretation
 FT
 FT XX
 FT PS Claim 11; Fig 4; 31pp; English.
 FT
 FT XX This sequence represents the upstream sequence, positions -611 to 9
 FT in the patent, of uridine diphosphate glucuronosyltransferase (UGT)
 FT gene 1*1 exon 1. This gene is known to be associated with Gilbert's
 FT syndrome (GS). GS is a mild, common form of unconjugated
 FT hyperbilirubinaemia associated with reduced bilirubin glucuronidation
 FT capacity. Analysis of the genetic basis of GS has allowed 2 forms to be
 FT identified. One is a mild form associated with a homozygous 2 bp
 FT insertion in the TATA sequence upstream of the UGT*1 exon 1, and
 FT the other is a more severe form associated with heterozygosity for a
 FT mutation which, when homozygous, causes Crigler-Najjar type 2 disease.
 FT The first form is autosomal recessive and the second is inherited
 FT dominantly. Patients suffering from GS, which is benign, may have
 FT altered metabolism of some drugs, making it difficult to determine if an
 FT effect is due to the drug or the syndrome. Drug trial efficiency would
 FT be improved if potential participants can be screened for the genetic
 FT basis of GS, and eliminated or included on basis of them possessing or
 FT not possessing GS. In this case, screening involves PCR amplification
 FT of the UGT gene, using the primers described in AAT79541-44.
 FT
 FT XX
 SQ Sequence 620 BP; 157 A; 127 C; 151 G; 185 T; 0 other:
 Query Match 100.0%; Score 21; DB 18; Length 620;
 Best Local Similarity 100.0%; Pred. No. 0.21;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 aagtaacctcctgctacctt 21
 DB 1 aagtaacctcctgctacctt 21

DB 465 aagtaactccctgctacctt 485

RESULT 3

AAV57903

ID AAV57903 standard; DNA; 237326 BP.

XX AAV57903;

DT 21-DEC-1998 (first entry)

DE Hereditary haemochromatosis subregion from an HH affected individual.

XX Bovine butyrophilin; BT; human hereditary haemochromatosis; HFE;

KW diagnosis; iron metabolism; NPT3; NPT4; Roret; BTF1; BTF2; BTF3;

KW BTF4; BTF5; milk protein; lupus; Sjogren's syndrome; hypophosphatemia;

XX type 1 sodium transport gene; ss.

XX Homo sapiens.

XX W09814466-A1.

XX 09-APR-1998.

XX 30-SEP-1997; 97MO-US17658.

XX 07-MAY-1997; 97US-0852495.

XX 01-OCT-1996; 96US-0724394.

XX (PROG-) PROGENITOR INC.

XX Feder JN, Krommal GS, Lauer PM, Ruddy DA, Thomas NJ;

XX Tsuchihashi Z, Wolff RK;

XX WPI; 1998-240014/21.

XX Hereditary haemochromatosis gene products - used to develop products

XX for the diagnosis and treatment of hereditary disorders in iron

XX metabolism

XX Claim 1; Fig 9; 209pp; English.

XX The present invention describes hereditary haemochromatosis gene

XX products from the human haemochromatosis gene. The present sequence

XX represents a hereditary haemochromatosis subregion from an hereditary

XX haemochromatosis (HH) affected individual. Also described is a

XX method to determine the presence or absence of the common hereditary

XX haemochromatosis (HFE) gene mutation in an individual comprising the

XX (a) providing DNA or RNA from the individual; and (b) assessing the

XX DNA or RNA for the presence or absence of a haplotype or genotype where

XX the presence or absence of the haplotype genotype indicates the likely

XX presence of the HFE gene mutation in the genome of the individual. The

XX HFE gene sequences from the present invention can be used to develop

XX products for use in the diagnosis and treatment of HFE. The present

XX invention also describes BTF genes, which are homologues of the milk

XX protein butyrophilin (BTF), and can be used in the production of agonists

XX and antagonists of BTF function. Also described are: (1) a Roret gene

XX which can be used to develop products for the study, diagnosis and

XX treatment of lupus and Sjogren's syndrome; and (2) NPT3 and NPT4 genes

XX which are homologues of a type 1 sodium transport gene, and can

XX similarly be used for hypophosphatemia.

XX Sequence 237326 BP; 69596 A; 48904 C; 48217 G; 70609 T; 0 other:

Query Match 80.0%; Score 16.8; DB 19; Length 237326;

Best Local Similarity 90.0%; Pred. No. 53;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 aagtaactccctgctacctt 20

DB 105327 aagtaactccctgctacctt 105346

RESULT 4

AAC25716/C

ID AAC25716 standard; CDNA; 260 BP.

XX AAC25716;

DT 06-OCT-2000 (first entry)

DE Human secreted protein 5' EST, SEQ ID NO: 29791.

XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;

XX gene therapy; chromosome mapping; ss.

XX Homo sapiens.

XX EP1033401-A2.

XX 06-SEP-2000.

XX 21-FEB-2000; 2000EP-0200610.

XX 26-FEB-1999; 99US-0122487.

XX (GEST) GENSET.

XX Dumas Maline Edwards J, Duclet A, Giordano J;

XX WPI; 2000-500381/45.

XX Claim 1; SEQ ID 29791; 71pp + CD-ROM; English.

XX The present sequence is one of a large number of 5' ESTs derived from

XX mRNAs encoding secreted proteins. No ORF has yet been conclusively

XX identified within the present sequence. The 5' ESTs were prepared from

XX total human RNAs or poly(A) RNAs derived from 30 different tissues. EST

XX sequences usually correspond mainly to the 3' untranslated region (UTR)

XX of the mRNA because they are often obtained from oligo-dT primed cDNA

XX libraries. Such ESTs are not well suited for isolating cDNA sequences

XX derived from the 5' ends of mRNAs and even in those cases where longer

XX cDNA sequences have been obtained, the full 5' UTR is rarely included.

XX 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be

XX used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used

XX in diagnostic, forensic, gene therapy and chromosome mapping procedures.

XX They are used to obtain upstream regulatory sequences and to design

XX expression and secretion vectors.

XX Sequence 260 BP; 74 A; 58 C; 45 G; 82 T; 1 other:

Query Match 77.1%; Score 16.2; DB 21; Length 260;

Best Local Similarity 85.7%; Pred. No. 46;

Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 aagtaactccctgctacctt 21

DB 68 AAGGAACCTGCTCTACCTT 48

RESULT 5

AAC38612/C

ID AAC38612 standard; DNA; 867 BP.

XX AAC38612;

DT 17-OCT-2000 (first entry)

DE Arabidopsis thaliana DNA fragment SEQ ID NO: 21606.

XX

Hybridisation assay; genetic mapping; gene expression control;
Protein identification; signal transduction pathway;
Metabolic pathway; promoter; termination sequence; ss.
Arabidopsis thaliana.
EPI033405-A2.
06-SEP-2000.
25-FEB-2000; 2000EP-0301439.
25-FEB-1999; 99US-0121825.
05-MAR-1999; 99US-0123180.
09-MAR-1999; 99US-0123548.
23-MAR-1999; 99US-0125788.
25-MAR-1999; 99US-0126264.
01-APR-1999; 99US-0127462.
06-APR-1999; 99US-0128234.
08-APR-1999; 99US-0128714.
15-APR-1999; 99US-0129845.
21-APR-1999; 99US-0130077.
23-APR-1999; 99US-0130449.
28-APR-1999; 99US-0130510.
30-APR-1999; 99US-0132048.
04-MAY-1999; 99US-0132407.
05-MAY-1999; 99US-0132484.
06-MAY-1999; 99US-0132485.
06-MAY-1999; 99US-0132486.
07-MAY-1999; 99US-0132863.
11-MAY-1999; 99US-0134256.
14-MAY-1999; 99US-0134218.
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18-MAY-1999; 99US-0134370.
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20-MAY-1999; 99US-0134941.
21-MAY-1999; 99US-0135124.
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24-MAY-1999; 99US-0135629.
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27-MAY-1999; 99US-0136392.
28-MAY-1999; 99US-0136582.
01-JUN-1999; 99US-0137222.
03-JUN-1999; 99US-0137528.
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18-JUN-1999; 99US-0139462.
18-JUN-1999; 99US-0139463.
18-JUN-1999; 99US-0139750.
18-JUN-1999; 99US-0139763.
21-JUN-1999; 99US-0139817.
22-JUN-1999; 99US-0139899.
23-JUN-1999; 99US-0140353.

23-JUN-1999; 99US-0140354.
24-JUN-1999; 99US-0140695.
28-JUN-1999; 99US-0140823.
29-JUN-1999; 99US-0140991.
30-JUN-1999; 99US-0141287.
01-JUL-1999; 99US-0141682.
01-JUL-1999; 99US-0142154.
02-JUL-1999; 99US-0142055.
06-JUL-1999; 99US-0142390.
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09-JUL-1999; 99US-0142920.
12-JUL-1999; 99US-0142977.
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27-JUL-1999; 99US-0145919.
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02-AUG-1999; 99US-0146386.
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17-AUG-1999; 99US-0149175.
18-AUG-1999; 99US-0149426.
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20-AUG-1999; 99US-0149922.
23-AUG-1999; 99US-0149902.
23-AUG-1999; 99US-0149930.
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27-AUG-1999; 99US-0151066.
27-AUG-1999; 99US-0151080.
30-AUG-1999; 99US-0151303.
31-AUG-1999; 99US-0151438.

PR 01-SEP-1999; 99US-0151930.
 PR 07-SEP-1999; 99US-0152363.
 PR 10-SEP-1999; 99US-0153070.
 PR 13-SEP-1999; 99US-0153758.
 PR 15-SEP-1999; 99US-0154018.
 PR 16-SEP-1999; 99US-0154039.
 PR 20-SEP-1999; 99US-0154779.
 PR 22-SEP-1999; 99US-0155139.
 PR 23-SEP-1999; 99US-0155486.
 PR 24-SEP-1999; 99US-0155659.
 PR 28-SEP-1999; 99US-0156458.
 PR 29-SEP-1999; 99US-0156596.
 PR 04-OCT-1999; 99US-0157117.
 PR 05-OCT-1999; 99US-0157573.
 PR 06-OCT-1999; 99US-0157865.
 PR 07-OCT-1999; 99US-0158029.
 PR 08-OCT-1999; 99US-0158232.
 PR 12-OCT-1999; 99US-0158369.
 PR 13-OCT-1999; 99US-0159293.
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 PR 14-OCT-1999; 99US-0159329.
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 PR 18-OCT-1999; 99US-0159584.
 PR 21-OCT-1999; 99US-0160741.
 PR 21-OCT-1999; 99US-0160767.
 PR 21-OCT-1999; 99US-0160770.
 PR 21-OCT-1999; 99US-0160814.
 PR 21-OCT-1999; 99US-0160815.
 PR 22-OCT-1999; 99US-0160980.
 PR 22-OCT-1999; 99US-0160981.
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 PR 25-OCT-1999; 99US-0161404.
 PR 25-OCT-1999; 99US-0161405.
 PR 25-OCT-1999; 99US-0161406.
 PR 26-OCT-1999; 99US-0161359.
 PR 26-OCT-1999; 99US-0161360.
 PR 26-OCT-1999; 99US-0161361.
 PR 28-OCT-1999; 99US-0161920.
 PR 28-OCT-1999; 99US-0161992.
 PR 28-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match 77.1%; Score 16.2; DB 21; Length 867;
 Best Local Similarity 85.7%; Pred. No. 53;
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 aagtgaaactccctgtaacct 21
 |||||
 DB 678 AAGTGAACCTCTTCGACACTT 658

RESULT 6
 AAA46500/c
 ID AAA46500 standard; cDNA; 1613 BP.
 XX
 AC AAA46500;
 DT 04-SEP-2000 (first entry)
 DE cDNA encoding enzyme involved in mannan polysaccharide hydrolysis.
 XX
 KW Hydrolysis; polysaccharide; mannan; coffee; endo-beta-mannanase; ds.
 XX
 OS Coffea arabica.
 XX
 FH Key 11.1294 Location/Qualifiers
 FT CDS /tag- a
 FT

FT /product- "endo-beta-mannanase"
 PN MO200028046-A1.
 PD 18-MAY-2000.
 PE 28-OCT-1999; 99MO-EP08314.
 PR 11-NOV-1999; 98EP-0203742.
 PA (NEST) SOC PROD NESTLE SA.
 PI Marzocchi P, Rogers J;
 DR WPI; 2000-399535/34.
 DR P-PSDB; AA93441.
 PT New DNA encoding endo-beta-mannanase from coffee, used e.g. in
 PT pharmaceutical, cosmetic or food compositions to hydrolyze polymannans
 PT
 PS Claim 4; Page 29-30; 41pp; French.

The present sequence encodes an endo-beta-mannanase enzyme, which is involved in the hydrolysis of polysaccharides that consist of molecules of mannan, either simple or branched, linked together by beta(1-4) bonds. The mannanase polynucleotide sequence is used for in vivo modification of the coffee endo-beta-mannanase gene. It is also used to produce transgenic plant cells (especially coffee cells) which have modified structures of mannan polysaccharide, and thus altered flavour or of mannan polysaccharides in vitro, particularly to treat coffee beans to increase the percentage of dry matter extraction, and thus reduce the quantity of sediment.

Sequence 1613 BP; 520 A; 305 C; 347 G; 441 T; 0 other;

Query Match 77.1%; Score 16.2; DB 21; Length 1613;
 Best Local Similarity 85.7%; Pred. No. 58;
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 aagtgaaactccctgtaacct 21
 |||||
 DB 408 AAGTGAACCTCTTCGACACTT 388

RESULT 7
 AA209252
 ID AA209252 standard; DNA; 32042 BP.
 XX
 AC AA209252;
 DT 25-OCT-1999 (first entry)
 DE Human CARD-4 DNA.

CARD-3; caspase recruitment domain; CARD-4; regulation; detection; caspase activation; detection; screening; therapy; diagnosis; disease; apoptotic cell death; Fas/Apo-1 receptor complex; TNF receptor complex; cancer; follicular lymphoma; carcinoma; p53 mutation; viral infection; hormone-dependent tumour; autoimmune disorder; Alzheimer's disease; systemic lupus erythematosus; immune-mediated glomerulonephritis; stroke; Parkinson's disease; amyotrophic lateral sclerosis; retinitis pigmentosa; spinal muscular dystrophy; cerebellar degeneration; anaemia; drug; myelodysplastic syndrome; myocardial infarction; cell proliferation; cell differentiation; cell survival; CARD-4L; CARD-4S; CARD-4Y; CARD-4Z; human; ds.

OS Homo sapiens.
 XX
 FH Key 364.685 Location/Qualifiers
 FT exon
 FT

PR	06-FEB-1998;	98US-0019942.
PR	PM 17-JUN-1998;	98US-0099041.
XX	(MILL-) MILLENNITUM PHARM INC.	
XX	Bertin J;	
DR	WPI; 1999-49A269/A1.	
XX		
PT	Novel CARD-3 and CARD-4 genes and polypeptides used or treating	
PT	regulation of cellular proliferation and differentiation and cell	
PT	survival	
PS	Example 13; Fig 18; 181pp; English.	
XX		
CC	This invention describes the isolation of novel human caspase	
CC	recruitment domain, CARD-3 and CARD-4 polynucleotides and proteins and a	
CC	partial murine CARD-4 protein and genes. The genes and proteins of	
CC	the invention are involved in the regulation of caspase activation.	
CC	The caspase recruitment domain (CARD) polynucleotides, polypeptides,	
CC	homologues and antibodies can be used in screening assays, detection	
CC	assays, predictive medicine and therapeutic and prophylactic methods of	
CC	treatment. The methods may be used to diagnose and treat patients which	
CC	are suffering from a disorder associated with abnormal level or rate of	
CC	apoptotic cell death, abnormal activity of the Fas/Apo-1 receptor	
CC	complex, abnormal activity of the TNF receptor complex, or abnormal	
CC	activity of a caspase. Diseases that may be treated include cancer	
CC	(particularly follicular lymphoma, carcinomas associated with mutations	
CC	in p53 and hormone-dependent tumours), autoimmune disorders (e.g.,	
CC	systemic lupus erythematosus, immune-mediated glomerulonephritis), viral	
CC	infections, Alzheimer's disease, Parkinson's disease, amyotrophic lateral	
CC	sclerosis, retinitis pigmentosa, spinal muscular dystrophy, cerebellar	
CC	dysgenesis, anaemia, myelodysplastic syndrome, myocardial infarction,	
CC	and stroke. CARD-3 protein interacts with other cellular proteins, and so	
CC	can be used for regulation of cellular proliferation and differentiation	
CC	or cell survival. The CARD proteins may also be used to for screen drugs	
CC	or compounds which modulate their activity. The CARD-4 gene can express a	
CC	long transcript that encodes CARD-4L, a short transcript that encodes	
CC	CARD-4S or two CARD-4 splice variants, CARD-4Y and CARD-4Z. This sequence	
CC	represents a genomic DNA sequence which encodes the human CARD-4 protein.	
XX		
SQ	Sequence 32042 BP; 7389 A; 7540 C; 7721 G; 9392 T; 0 other;	
OY	1 aaagtaactcctgccttcctt 21	
Dy	17935 aaaggaaactccctgatacctt 17955	
RESULT 8		
AFAF30011		
ID AAF30011 standard; cDNA; 32042 BP.		
XX		
AC AAF30011:		
DT 23-APR-2001 (first entry)		
DE XX		
XX Human CARD-4 gene.		
KW CARD-4; caspase recruitment domain; human; cancer; infection;		
KW autoimmune disease; neurological disease; hematological disease;		
KW immune disease; inflammation; antitumour; antiapptic;		
KW immunomodulator; antiinflammatory; apoptosis; diagnosis;		
KW gene therapy; chromosome 7; ds.		
OS Homo sapiens.		
FH Key Location/Qualifiers		
FT CDS 485..31768		

```

FT      /*tag- a
FT      /note- "Contains Introns"
FT      PA      364..685
FT      PI      /*tag- b
FT      DR      /number- 1
FT      DR      686..2094
FT      DR      /*tag- c
FT      DR      /number- 1
FT      DR      2095..2269
FT      DR      /*tag- d
FT      DR      /number- 2
FT      DR      2270..4365
FT      DR      /*tag- e
FT      DR      /number- 2
FT      DR      4366..6190
FT      DR      /*tag- f
FT      DR      /number- 3
FT      DR      6191..9024
FT      DR      /*tag- g
FT      DR      /number- 3
FT      DR      9025..9108
FT      DR      /*tag- h
FT      DR      /number- 4
FT      DR      9109..10355
FT      DR      /*tag- i
FT      DR      /number- 4
FT      DR      10356..10439
FT      DR      /*tag- j
FT      DR      /number- 5
FT      DR      10440..11181
FT      DR      /*tag- k
FT      DR      /number- 5
FT      DR      11182..11265
FT      DR      /*tag- l
FT      DR      /number- 6
FT      DR      11266..19749
FT      DR      /*tag- m
FT      DR      /number- 6
FT      DR      19750..19833
FT      DR      /*tag- n
FT      DR      /number- 7
FT      DR      19834..21324
FT      DR      /*tag- o
FT      DR      /number- 7
FT      DR      21325..21408
FT      DR      /*tag- p
FT      DR      /number- 8
FT      DR      21409..24226
FT      DR      /*tag- q
FT      DR      /number- 8
FT      DR      24227..24310
FT      DR      /*tag- r
FT      DR      /number- 9
FT      DR      24311..27948
FT      DR      /*tag- s
FT      DR      /number- 9
FT      DR      27949..28032
FT      DR      /*tag- t
FT      DR      /number- 10
FT      DR      28033..31695
FT      DR      /*tag- u
FT      DR      /number- 10
FT      DR      31696..32024
FT      DR      /*tag- v
FT      DR      /number- 11

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XX      MO200100826-A2.
XX      PD      04-JAN-2001.
XX      PF      28-JUN-2000. 2000M0-DS17691.
XX      PR      28-JUN-1999. 9905-0340620.

```

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XX      (MILL-) MILLENNIUM PHARM INC.
XX      PI      Bertha J.
XX      DR      WPI: 2001-061973/07.
XX      DR      P-PSDB: AAB20080, AAB20081, AAB20082, AAB20083.
XX      PT      Isolated intracellular proteins predicted to be involved in regulating
XX      PT      caspase activation are used for diagnosis and treatment of e.g. cancer,
XX      PT      viral infections, autoimmune diseases, neurological diseases and
XX      PT      haematological disorders.
XX      PS      Example 13: Fig 18: 208pp; English.
XX      CC      The present sequence is that of the human caspase recruitment
XX      CC      domain 4 (CARD-4) gene on chromosome 7. CARD-4 exists in at least
XX      CC      4 forms, i.e. the long form CARD-4L (see AAB20080), the short form
XX      CC      CARD-4S (see AAB20081), and splice variants CARD-4Y (see AAB20082)
XX      CC      and CARD-4Z (see AAB20082). It is an intracellular protein
XX      CC      predicted to be involved in regulating caspase activation. It
XX      CC      activates the NF-kappaB pathway and enhances caspase-9-mediated
XX      CC      cell death. Methods of diagnosing and treating patients suffering
XX      CC      from a disorder associated with an abnormal level or rate of apoptotic
XX      CC      cell death, abnormal activity of the Fas/Apo-1 receptor complex,
XX      CC      abnormal activity of the tumour necrosis factor receptor complex,
XX      CC      or abnormal activity of a caspase involve administering a compound
XX      CC      that modulates the expression or activity of CARD-3, CARD-4, CARD-5
XX      CC      or CARD-6 e.g. a small molecule, antisense nucleic acid, ribozyme
XX      CC      or polypeptide. Such disorders include cancer, viral infection,
XX      CC      autoimmune disorders, neurological diseases, haematological
XX      CC      disorders, inflammatory disorders and immune disorders. The CARD-4
XX      CC      gene is useful for genetic information and mapping and identifying
XX      CC      mutations, e.g. mutations in splice donor or acceptor sites.
XX      SO      Sequence 32042 BP: 7389 A; 7540 C; 7721 G; 9392 T; 0 other;

Query Match      77.1%; Score 16.2; DB 22; Length 32042;
Best Local Similarity 85.7%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY      1 aagtgaaacctccctgacctt 21
DB      17935 aagsgaaacctccctgacctt 17955

RESULT 9
AA097865
ID      AA097865 standard; DNA; 957 BP.
XX      AC      AA097865;
XX      DT      05-DEC-1995 (first entry)
XX      DE      TROMP1 DNA.
XX      DE      TROMP1 DNA.
XX      KW      TROMP1; rare outer membrane protein; Spirochaetaceae; immunogen;
XX      KW      vaccine; syphilis; ss.
XX      OS      Treponema pallidum subsp. pallidum.
XX      FH      Key      Location/Qualifiers
XX      FT      CDS      1..957
XX      FT      sig_peptide 1..96
XX      FT      mat_peptide 97..954
XX      FT      /*tag- a
XX      FT      /*tag- b
XX      FT      /*tag- c
XX      PD      MO9518632-A.
XX      PD      13-JUL-1995.

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XX 06-JAN-1995; 95WO-US00190.
PF 17-AUG-1994; 94US-0292904.
XX 06-JAN-1994; 94US-0178084.
PR 07-JUN-1994; 94US-0255322.
XX (REGC ) UNIV CALIFORNIA.
XX Blanco DR, Champion CT, Lovett MA, Miller JN;
PI WPI; 1995-254907/33.
DR P-PSDB; AAR79722.
XX Rare outer coat membranes of Spirochaetaeaceae prodn. - by density
PT gradient fractionation, also isolated immunogenic proteins for use in
PT vaccines, derived antibodies and nucleic acid.
XX Disclosure; Page 48-50; 78pp; English.
PS TROMPI (given in AAR79722) is encoded by DNA isolated using tryptic
CC digest amino acid sequence analysis of EcoRI-digested genomic DNA
CC of T. pallidum subsp. pallidum. It is the precursor of a 31 kDa rare
CC outer membrane protein useful for prepn. of vaccine against syphilis
CC and other treponemal diseases.
XX Sequence 957 BP; 210 A; 206 C; 303 G; 238 T; 0 other;
SQ
Query Match 75.2%; Score 15.8; DB 16; Length 957;
Best Local Similarity 89.5%; Pred. No. 85;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY 3 gtaactccctgcatactt 21
DB 749 gtaactccctgcatactt 767
RESULT 10
AAT6580
ID AAT6580 standard; DNA; 957 BP.
XX AAT6580;
AC 13-JAN-1998 (first entry)
XX Treponema pallidum rare outer membrane protein (TROMP-1) DNA.
DE Treponema pallidum rare outer membrane protein (TROMP-1) DNA.
XX Rate: outer membrane protein; Treponema pallidum; Spirochaetales;
KM Immune response; Syphilis; TROMP-1; precursor; ss.
XX Treponema pallidum.
OS
XX Key Location/Qualifiers
FH 1..957
FT CDS /tag= a
FT /product= Immature_TROMP1_protein
FT
XX MO972725-A1.
XX 31-JUL-1997.
XX 23-JAN-1997; 97WO-US01302.
XX 23-JAN-1996; 96US-0599480.
XX 23-JAN-1996; 96US-0599480.
XX (SLOK ) SLOAN KETTERING INST CANCER RES.
XX (REGC ) UNIV CALIFORNIA.
XX Blanco DR, Lovitt MA, Miller JN, Tempst PJ;
PI WPI; 1997-393614/36.
DR P-PSDB; AAR22134.

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XX New isolated Treponema pallidum outer membrane protein - used in the
PT detection and induction of immune response in an animal to
PT pathogenic Spirochaetales
XX Disclosure; Pages 51-53; 77pp; English.
PS This 957 bp sequence (TROMPI gene) encodes a precursor rare outer
CC membrane protein (OMP) of species Treponema pallidum, subspecies
CC pallidum. The TROMPI gene was cloned in a procedure where mixed
CC oligonucleotides (31-A, 31-C) hybridized to a EcoRI restriction
CC fragment by Southern blot analysis of T. pallidum genomic DNA. DNA
CC fragments were excised from the agarose gel, purified and ligated
CC into the lambda Zap II vector and probed with 31-A and 31-C. The
CC phage clones were converted to the Bluescript SK(-) recombinant
CC plasmid by in vivo excision. The recombinant expression of these
CC rare OMP's can be used for diagnostic tests to detect syphilis and
CC for development of host immunity during syphilis.
XX Sequence 957 BP; 210 A; 206 C; 303 G; 238 T; 0 other;
SQ
Query Match 75.2%; Score 15.8; DB 18; Length 957;
Best Local Similarity 89.5%; Pred. No. 85;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY 3 gtaactccctgcatactt 21
DB 749 gtaactccctgcatactt 767
RESULT 11
AAA67251/C
ID AAA67251 standard; DNA; 537 BP.
XX AAA67251;
AC 31-OCT-2000 (first entry)
XX Eucalyptus grandis beta-amylase DNA sequence SEQ ID NO:252.
DE Eucalyptus grandis beta-amylase DNA sequence SEQ ID NO:252.
XX Eucalyptus grandis; Pinus radiata; Monterey pine; modification;
KM plant cell wall; polysaccharide; polysaccharide biosynthetic pathway;
KM transgenic plant; ds.
XX Eucalyptus grandis.
OS
XX WO200022092-A2.
XX 20-APR-2000.
XX 08-OCT-1999; 99WO-N200169.
XX 13-OCT-1998; 98US-0170862.
XX 11-AUG-1999; 99US-0148426.
XX (GENE-) GENESIS RES & DEV CORP LTD.
XX (FLET-) FLETCHER CHALLENGE FORESTS LTD.
XX Bloksberg LN;
XX WPI; 2000-339328/29.
XX New genes encoding proteins involved in a plant polysaccharide
PT biosynthetic pathway, useful for modulating or altering the
PT polysaccharide content, composition or structure of the plant -
XX Claim 1; Page 140; 301pp; English.
XX The present invention describes isolated polynucleotides (PN) comprising
CC a sequence selected from one of 835 nucleotide sequences given in
CC AAA67073 to AAA67907, their (reverse) complements, sequences producing
CC an Expectation (E) value of 0.01 or less compared to the 835 sequences,

```

CC sequences at least 50% identical to them, 200, 100, 40 or 20-mers of the
 CC 835 sequences or sequences that are degenerately equivalent or allelic
 CC to the 835 sequences. The polynucleotides are used to modify the
 CC activity of a polypeptide involved in a polysaccharide biosynthetic
 CC pathway in the plant. They are especially used to modulate or alter the
 CC polysaccharide content, composition or structure of the plant. AAB16268
 CC to AAB16340 are proteins encoded by some of the polynucleotide sequence
 CC given in the present invention.

Sequence 537 BP; 155 A; 119 G; 136 C; 127 T; 0 other;

Query Match 72.4%; Score 15.2; DB 21; Length 537;
 Best Local Similarity 85.0%; Pred. No. 1.6e+02;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 agtgaactccctgctactct 21
 ||||| ||||| |||||
 Db 472 AGTGAATCTCTGCAACCTT 453

RESULT 12

ID AAV04426 standard; CDNA; 640 BP.

AC AAV04426;

DT 27-APR-1998 (first entry)

DE Human calpain small subunit cDNA (fragment).

XX Calpain; human; leukocyte; calcium dependent cysteine protease;
 KW screening; activator; inhibitor; treatment; prevention; cancer;
 KW cerebral apoplexy; cerebral infarction; subarachnoid haemorrhage;
 KW Alzheimer's disease; myodystrophy; cataracts; collagen disease;
 KW ischemic heart disease; atherosclerosis; arthritis;
 KW small subunit; ds.

XX Homo sapiens.

XX EP799892-A2.

XX 08-OCT-1997.

XX 03-APR-1997; 97EP-0105508.

XX 05-APR-1996; 96JP-0083649.

XX (TAKE) TAKEDA CHEM IND LTD.

XX Kawamoto T, Nishi K, Shintani Y;

XX WPI: 1997-482674/45.

XX Human calpain protein and related DNA - useful for drug screening
 PT and treating cancer, stroke, etc.

XX Example 2; Page 38; 43pp; English.

XX The present sequence is a calpain small subunit cDNA fragment.
 CC Calpain is a human leukocyte derived calcium dependent cysteine
 CC protease. Calpain can be used to screen for compounds that activate
 CC or inhibit its proteolytic activity. Calpain DNA can be used to
 CC treat or prevent cancer, cerebral apoplexy, cerebral infarction,
 CC subarachnoid haemorrhage, Alzheimer's disease, myodystrophy,
 CC cataracts, ischemic heart disease, atherosclerosis, arthritis
 CC or collagen disease.

Sequence 640 BP; 139 A; 163 C; 223 G; 115 T; 0 other;

Query Match

Best Local Similarity 85.0%; Pred. No. 1.6e+02;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 agtgaactccctgctactct 21
 ||||| ||||| |||||
 Db 574 AGTGAATCTCTGCAACCTT 593

RESULT 13

ID AAV04420 standard; CDNA; 804 BP.

AC AAV04420;

DT 27-APR-1998 (first entry)

DE Human calpain small subunit cDNA.

XX Calpain; human; leukocyte; calcium dependent cysteine protease;
 KW screening; activator; inhibitor; treatment; prevention; cancer;
 KW cerebral apoplexy; cerebral infarction; subarachnoid haemorrhage;
 KW Alzheimer's disease; myodystrophy; cataracts; collagen disease;
 KW ischemic heart disease; atherosclerosis; arthritis;
 KW small subunit; ds.

XX Homo sapiens.

XX EP799892-A2.

XX 08-OCT-1997.

XX 03-APR-1997; 97EP-0105508.

XX 05-APR-1996; 96JP-0083649.

XX (TAKE) TAKEDA CHEM IND LTD.

XX Kawamoto T, Nishi K, Shintani Y;

XX WPI: 1997-482674/45.

XX Human calpain protein and related DNA - useful for drug screening
 PT and treating cancer, stroke, etc.

XX Example 2; Page 35; 43pp; English.

XX The present sequence encodes calpain small subunit, a human
 CC leukocyte derived calcium dependent cysteine protease. Calpain can
 CC be used to screen for compounds that activate or inhibit its
 CC proteolytic activity. Calpain DNA can be used to treat or prevent
 CC cancer, cerebral apoplexy, cerebral infarction, subarachnoid
 CC haemorrhage, Alzheimer's disease, myodystrophy, cataracts,
 CC ischemic heart disease, atherosclerosis, arthritis or collagen
 CC disease.

Sequence 804 BP; 181 A; 203 C; 264 G; 156 T; 0 other;

Query Match

Best Local Similarity 85.0%; Pred. No. 1.7e+02;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 agtgaactccctgctactct 21
 ||||| ||||| |||||
 Db 574 AGTGAATCTCTGCAACCTT 593

RESULT 14

AC7765
ID AAC7765 standard; cDNA, 1242 BP.
AC AAC7765;
DT 08-FEB-2001 (first entry)
DE Human cancer associated gene sequence SEQ ID NO:159.
XX
XX Human; cancer associated gene; cancer antigen; detection; cancer;
XX diagnosis; cytostatic; proliferative; vulnery; immunomodulator;
XX antidiabetic; antihypertensive; antirheumatic; antitumor; antiviral;
XX antiinflammatory; antihypertensive; antitumor; antidiabetic; antitumor;
XX dermatological; neuroprotective; thrombolytic; coagulant; neutropenic;
XX vasotropic; antiproliferative; antitumor; gene therapy; inflammation;
XX immune disorder; hematopoietic cell disorder; autoimmune disorder;
XX allergic reaction; graft versus host disease; organ rejection;
XX haemostatic; thrombolytic; cardiovascular disorder; infection;
XX neurological disease; drug screening; ss.
OS Homo sapiens.
PN WO20005350-A1.
PD 21-SEP-2000.
PF 08-MAR-2000; 2000WO-US05882.
PR 12-MAR-1999; 99US-0124270.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM;
XX WPI: 2000-587533/55.
DR P-PSDB: ABA43556.
DR
XX Novel isolated nucleic acids comprising sequences encoding peptides
PT useful for treating or diagnosing e.g. cancer -
XX
XX Claim 1: Page 737-738; 2352pp; English.
PS
CC AAC77657 to AAC77658 encode the human cancer associated proteins given
CC in AAB3398 to AAB4423. The proteins can have activities based on the
CC tissues and cells the genes are expressed in. Example of activities
CC include: cytostatic; proliferative; vulnery; immunomodulator;
CC antidiabetic; antihypertensive; antirheumatic; antitumor; antiviral;
CC antiinflammatory; antihypertensive; antitumor; antidiabetic; antitumor;
CC dermatological; neuroprotective; thrombolytic; coagulant;
CC neutropenic; vasotropic; antiproliferative; antitumor; gene therapy;
CC inflammation; immune disorder; hematopoietic cell disorder; autoimmune disorder;
CC allergic reaction; graft versus host disease; organ rejection;
CC haemostatic; thrombolytic; cardiovascular disorder; infection;
CC neurological disease; drug screening; ss.
CC disorders, allergic reactions, graft versus host disease and organ
CC rejection, modulate haemostatic or thrombolytic activity, modulate
CC inflammation, cancers, cardiovascular disorders, neurological disease and
CC bacterial or viral infections. The peptides, nucleotides, antibodies,
CC agonists and antagonists may be also be used in drug screens. AAC77649 to
CC AAC77657 and ABA44240 represent sequences used in the exemplification of
CC the present invention.
XX
XX Sequence 1242 BP; 333 A; 357 C; 292 G; 258 T; 2 other;
XX
XX Query Match 72.4%; Score 15.2; DB 21; Length 1242;
XX Best local similarity 85.0%; Pred. No. 1.8e-02;
XX Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
XX 2 agtgaactccctcactcctt 21
XX ||||||||| | |||||

DB 414 agtgaactccctcactcctt 433
RESULT 15
AAZ98118/C
ID AAZ98118 standard; cDNA; 1741 BP.
XX
XX AAZ98118;
XX
DT 11-MAY-2000 (first entry)
DE Human signal peptide containing protein HSP-10 cDNA SEQ ID NO:144.
XX
XX Human; signal peptide-containing protein; HSP; diagnosis; cancer;
XX inflammation; cardiovascular disease; anticancer; anti-inflammatory;
XX antitumor; neuroprotective; cardiovascular; hepatotropic;
XX antidiabetic; gene therapy; cell proliferation; neurological disorder;
XX antitumor; antihypertensive; antitumor; antidiabetic; antitumor;
XX reproductive disorder; developmental disorder; arteriosclerosis;
XX cirrhosis; psoriasis; acquired immune deficiency syndrome; anaemia;
XX asthma; Crohn's disease; infection; Alzheimer's disease; schizophrenia;
XX Parkinson's disease; Huntington's disease; ovulatory defect;
XX muscular dystrophy; ss.
OS Homo sapiens.
PN WO20000610-A2.
PD 06-JAN-2000.
PF 25-JUN-1999; 99WO-US14484.
PR 26-JUN-1998; 98US-0090762.
PR 31-JUL-1998; 98US-0094983.
PR 01-OCT-1998; 98US-0102686.
PR 11-DEC-1998; 98US-0112129.
XX
XX (INCY-) INCYTE PHARM INC.
XX
XX Lal P, Tang YT, Gorgone GA, Corley NC, Guegler KI, Baughn MR;
XX Akeldom IE, Au-Young J, Yue H, Patterson C, Reddy R, Hillman JL;
XX Bandman O;
XX WPI: 2000-150673/14.
DR P-PSDB: AAY7233.
DR
XX New human signal peptide-containing proteins useful in treatment,
PT prevention and diagnosis of e.g. cancer, inflammation and
PT cardiovascular disease -
XX
XX Claim 9: Page 257-258; 327pp; English.
PS
CC AAZ98109 to AAZ98242 encode AAY7224 to AAY7357 which represent the
CC human signal peptide-containing proteins HSP-1 to HSP-134. HSPs have
CC anticancer, anti-inflammatory, antimicrobial, neutropenic, hepatotropic,
CC neuroprotective, cardiovascular and antitumor activities, and can
CC be used in gene therapy. HSPs can be used to treat or prevent disorders
CC associated with decreased activity or function of HSP. Antagonists of
CC HSP are used to treat or prevent disorders associated with increased
CC activity or function of HSP. Such diseases include cell proliferation
CC (including cancer), inflammation, cardiovascular, neurological,
CC reproductive or developmental disorders, (e.g. arteriosclerosis,
CC cirrhosis, psoriasis, acquired immune deficiency syndrome, anaemia,
CC asthma, Crohn's disease, microbial or other infections, congestive or
CC ischemic heart disease, Alzheimer's, Parkinson's or Huntington's
CC diseases, schizophrenia, ovulatory defects, muscular dystrophy). HSP
CC nucleic acids can be used for the recombinant production of HSP, for
CC detecting HSP in standard hybridisation and amplification assays (for
CC diagnosis and monitoring), in gene therapy, as antisense,
CC triplex-forming or ribozyme therapeutics, for detecting related sequences
CC or genetic variations, and for chromosomal mapping. HSP are also used to
CC raise specific antibodies (Ab) and to screen for agonists and
CC antagonists (potential therapeutic agents). Ab are used to diagnose, or
CC monitor, HSP-related diseases (in usual immunoassays), as therapeutic

CC antagonists, in competitive drug screens, and for purification of NSP
CC from natural sources.

XX
SQ Sequence 1741 BP; 533 A; 358 C; 364 G; 486 T; 0 other;

Query Match 72.4%; Score 15.2; DB 21; Length 1741;

Best Local Similarity 85.0%; Pred. No. 1.8e+02;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 aagtgactccgcctact 20

||||| | |||||

Db 1411 AAGTGAGTGTCTACT 1392

Search completed: July 25, 2001, 05:23:03
Job Time: 4676 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 02:45:17 ; Search time 117.39 seconds
(without alignments)
33.141 Million cell updates/sec

File: US-09-142-095-1

Perfect score: 21

Sequence: 1 aaagtaactccctctactcct 21

Scoring table:

IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 317530 seqs, 92630169 residues

Total number of hits satisfying chosen parameters: 635060

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: /cgn2_6/ptodata/2/1na/5A.COMB.seq:*
2: /cgn2_6/ptodata/2/1na/5B.COMB.seq:*
3: /cgn2_6/ptodata/2/1na/5A.COMB.seq:*
4: /cgn2_6/ptodata/2/1na/5B.COMB.seq:*
5: /cgn2_6/ptodata/2/1na/PCITUS.COMB.seq:*
6: /cgn2_6/ptodata/2/1na/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	15.8	75.2	957 1 US-08-599-480-1	Sequence 1, Appl
2	15.8	75.2	957 1 US-08-842-198-1	Sequence 1, Appl
3	15.8	75.2	957 5 PCT-US85-00180-1	Sequence 1, Appl
4	15.2	72.4	640 2 US-08-835-099A-16	Sequence 16, Appl
5	15.2	72.4	640 3 US-09-157-349-16	Sequence 16, Appl
6	15.2	72.4	804 2 US-08-835-099A-10	Sequence 10, Appl
7	15.2	72.4	804 3 US-09-157-349-10	Sequence 10, Appl
8	15.2	72.4	246240 2 US-08-724-394A-20	Sequence 20, Appl
9	15.2	72.4	246240 2 US-08-724-394A-20	Sequence 20, Appl
10	15.2	72.4	246240 2 US-08-724-394A-22	Sequence 22, Appl
11	14.8	70.5	11723 1 US-08-417-492-1	Sequence 1, Appl
12	14.8	70.5	2161 2 US-09-001-826-4	Sequence 1, Appl
13	14.8	70.5	2161 2 US-09-001-826-25	Sequence 25, Appl
14	14.8	70.5	2652 2 US-08-366-547-1	Sequence 1, Appl
15	14.8	70.5	2977 1 US-08-276-151-1	Sequence 1, Appl
16	14.8	70.5	2977 1 US-08-306-691B-17	Sequence 17, Appl
17	14.8	70.5	2977 1 US-08-756-806A-64	Sequence 64, Appl
18	14.8	70.5	2977 3 US-08-328-239A-4	Sequence 4, Appl
19	14.8	70.5	2977 3 US-09-143-214-64	Sequence 64, Appl
20	14.8	70.5	2977 3 US-09-209-668-12	Sequence 64, Appl
21	14.8	70.5	2977 5 PCT-US95-13661-4	Sequence 14, Appl
22	14.6	69.5	394 4 US-08-488-144-18	Sequence 18, Appl
23	14.6	69.5	1168 4 US-08-930-285-15	Sequence 15, Appl
24	14.6	69.5	4345 3 US-08-244-537-1	Sequence 5, Appl
25	14.6	69.5	7470 3 US-08-417-089-5	Sequence 5, Appl
26	14.6	69.5	7470 4 US-08-695-651-5	Sequence 5, Appl
27	14.6	69.5	7470 4 US-08-930-285-5	Sequence 5, Appl

28	14.4	68.6	1273 4 US-08-725-758A-3	Sequence 3, Appl
29	14.4	68.6	1373 4 US-08-725-758A-1	Sequence 1, Appl
30	14.4	68.6	4310 4 US-09-008-172-1	Sequence 1, Appl
31	14.2	67.6	61 4 US-08-687-421-355	Sequence 355, Appl
32	14.2	67.6	1118 1 US-08-418-032-1	Sequence 1, Appl
33	14.2	67.6	1554 1 US-08-469-486-1	Sequence 1, Appl
34	14.2	67.6	1554 2 US-08-469-486-1	Sequence 1, Appl
35	14.2	67.6	3504 1 US-08-620-717A-8	Sequence 8, Appl
36	14.2	67.6	5541 1 US-08-920-812-20	Sequence 20, Appl
37	14.2	67.6	5541 1 US-08-920-827-20	Sequence 20, Appl
38	14.2	67.6	5541 1 US-08-920-827-20	Sequence 20, Appl
39	14.2	67.6	5541 1 US-08-920-827-20	Sequence 20, Appl
40	14.2	67.6	5541 1 US-08-362-577C-20	Sequence 20, Appl
41	14.2	67.6	5541 2 US-08-920-828-20	Sequence 20, Appl
42	14.2	67.6	20303 1 US-08-370-975B-6	Sequence 6, Appl
43	14.2	67.6	26764 1 US-08-370-975B-1	Sequence 1, Appl
44	14.2	67.6	1441 1 US-08-136-277-18	Sequence 18, Appl
45	14.2	67.6	1441 2 US-08-479-403-18	Sequence 18, Appl
			1441 3 US-08-835-734-18	Sequence 18, Appl

ALIGNMENTS

RESULT 1
US-08-599-480-1
Sequence 1, Application US/08599480
Patent No. 5753459
GENERAL INFORMATION:
APPLICANT: Blanco, David R.
APPLICANT: Miller, James N.
APPLICANT: Loyett, Michael A.
APPLICANT: Champion, Cheryl I.
APPLICANT: Tempst, Paul J.
TITLE OF INVENTION: NUCLEOTIDE AND AMINO ACID SEQUENCES OF A
NUMBER OF SEQUENCES: 4
TITLE OF INVENTION: T. Pallidum RARE OUTER MEMBRANE PROTEIN
CORRESPONDENCE ADDRESS:
ADDRESS: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: California
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/599,480
FILING DATE: 23-JAN-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Learn, June M.
REGISTRATION NUMBER: 31,238
TELEPHONE/DOCKET NUMBER: 07419/018001 (CIP of 016001)
TELEPHONE: (619) 678-5070
TELEFAX: (619) 678-5099
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 957 base pairs
TYPE: nucleic acid
STANDARDS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
IMMEDIATE SOURCE:
CLONE: TROMP1
FEATURES:
NAME/KEY: CDS
LOCATION: 1..954
US-08-599-480-1

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,099A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 083649/1996
FILING DATE: 05-APR-1996
APPLICATION NUMBER: 97105508.2
FILING DATE: 03-APR-1997
ATTORNEY/AGENT INFORMATION:
NAME: Resnick, David S
REGISTRATION NUMBER: 34,235
REFERENCE/DOCKET NUMBER: 47342
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-523-3400
TELEFAX: 617-523-6440
TELEX:
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 640 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-08-835-099A-16

Query Match 72.4% Score 15.2; DB 2; Length 640;
Best Local Similarity 85.0%; Pred. No. 36;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 agtgaactccctgctacct 21
|||||
Db 574 agtgaactccctgctacct 593

RESULT 5
US-09-157-349-16
Sequence 16, Application US/09157349
Patent No. 6068990
GENERAL INFORMATION:
APPLICANT: SHINTANI, Yasushi
APPLICANT: NISHI, Kazuo
APPLICANT: KAWAMOTO, Tomohiro
TITLE OF INVENTION: NOVEL PROTEINS, THEIR PRODUCTION
TITLE OF INVENTION: AND USE
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: DIKE, BRONSTEIN, ROBERTS & CUSMAN, LLP
STREET: 130 Water Street
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02109
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/157,349
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/835,099
FILING DATE:
APPLICATION NUMBER: 97105508.2
FILING DATE: 03-APR-1997
ATTORNEY/AGENT INFORMATION:

NAME: Resnick, David S
REGISTRATION NUMBER: 34,235
REFERENCE/DOCKET NUMBER: 47342
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-523-3400
TELEFAX: 617-523-6440
TELEX:
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 640 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-09-157-349-16

Query Match 72.4% Score 15.2; DB 3; Length 640;
Best Local Similarity 85.0%; Pred. No. 36;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 agtgaactccctgctacct 21
|||||
Db 574 agtgaactccctgctacct 593

RESULT 6
US-08-835-099A-10
Sequence 10, Application US/08835099A
Patent No. 5874277
GENERAL INFORMATION:
APPLICANT: SHINTANI, Yasushi
APPLICANT: NISHI, Kazuo
APPLICANT: KAWAMOTO, Tomohiro
TITLE OF INVENTION: NOVEL PROTEINS, THEIR PRODUCTION
TITLE OF INVENTION: AND USE
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: DIKE, BRONSTEIN, ROBERTS & CUSMAN, LLP
STREET: 130 Water Street
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02109
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,099A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 083649/1996
FILING DATE: 05-APR-1996
APPLICATION NUMBER: 97105508.2
FILING DATE: 03-APR-1997
ATTORNEY/AGENT INFORMATION:
NAME: Resnick, David S
REGISTRATION NUMBER: 34,235
REFERENCE/DOCKET NUMBER: 47342
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-523-3400
TELEFAX: 617-523-6440
TELEX:
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 804 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA

US-08-835-099A-10

Query Match 72.4%; Score 15.2; DB 2; Length 804;

Best Local Similarity 85.0%; Pred. No. 37;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 agtgaactccctgctaccct 21
|||||

DB 574 AGTGACTCCAGGTGCTT 593

RESULT 7

US-09-157-349-10

Sequence 10, Application US/09157349

Patent No. 6068990

GENERAL INFORMATION:

APPLICANT: SHINTANI, Yasushi

APPLICANT: NISHI, Kazuo

APPLICANT: KAMAMOTO, Tomohito

TITLE OF INVENTION: NOVEL PROTEINS, THEIR PRODUCTION

NUMBER OF SEQUENCES: 18

CORRESPONDENCE ADDRESS:

ADDRESSEE: DIKE, BRONSTEIN, ROBERTS & CUSMAN, LLP

STREET: 130 Water Street

CITY: Boston

STATE: MA

COUNTRY: USA

ZIP: 02109

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

OPERATING SYSTEM: IBM Compatible

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/157,349

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/835,099

FILING DATE:

APPLICATION NUMBER: 97105508.2

FILING DATE: 03-APR-1997

ATTORNEY/AGENT INFORMATION:

NAME: Resnick, David S

REGISTRATION NUMBER: 34,235

REFERENCE/DOCKET NUMBER: 47342

TELECOMMUNICATION INFORMATION:

TELEPHONE: 617-523-3400

TELEFAX: 617-523-6440

TELEX:

INFORMATION FOR SEQ ID NO: 10:

SEQUENCE CHARACTERISTICS:

LENGTH: 804 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

MOLECULE TYPE: CDNA

US-09-157-349-10

Query Match 72.4%; Score 15.2; DB 3; Length 804;
Best Local Similarity 85.0%; Pred. No. 37;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 2 agtgaactccctgctaccct 21
|||||
DB 574 AGTGACTCCAGGTGCTT 593. RESULT 8
US-08-724-394A-20/C

Sequence 20, Application US/08724394A

Patent No. 5872237

GENERAL INFORMATION:

APPLICANT: Feder, John N.

APPLICANT: Kronmal, Gregory S.

APPLICANT: Lauer, Peter M.

APPLICANT: Ruddy, David A.

APPLICANT: Thomas, Winston

APPLICANT: Tsuchihashi, Zenta

APPLICANT: Wolff, Roger K.

TITLE OF INVENTION: Megabase Transcript Map: No. 5872237e1

NUMBER OF SEQUENCES: 31

CORRESPONDENCE ADDRESS:

ADDRESSEE: TOWNSEND and TOWNSEND and CREW LLP

STREET: Two Embarcadero Center, 8th Floor

CITY: San Francisco

STATE: CA

COUNTRY: USA

ZIP: 94111-3834

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/724,394A

FILING DATE: 01-OCT-1996

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Fitts, Renee A.

REGISTRATION NUMBER: 35,136

REFERENCE/DOCKET NUMBER: 017957-000100

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-576-0200

TELEFAX: 415-576-0300

INFORMATION FOR SEQ ID NO: 20:

SEQUENCE CHARACTERISTICS:

LENGTH: 246240 base pairs

TYPE: nucleic acid

STRANDEDNESS: not relevant

TOPOLOGY: not relevant

MOLECULE TYPE: CDNA

FEATURE:

NAME/KEY: misc_feature

LOCATION: 1..246240

OTHER INFORMATION: /note="HNA-H.CONTIG"

US-08-724-394A-20

Query Match 72.4%; Score 15.2; DB 2; Length 246240;
Best Local Similarity 85.0%; Pred. No. 86;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 aagtgaactccctgctaccct 20
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DB 139464 ATGTGACTCCTGCACTT 139445

RESULT 9

US-08-724-394A-21/C

Sequence 21, Application US/08724394A

Patent No. 5872237

GENERAL INFORMATION:

APPLICANT: Feder, John N.

APPLICANT: Kronmal, Gregory S.

APPLICANT: Lauer, Peter M.

APPLICANT: Ruddy, David A.

APPLICANT: Thomas, Winston

APPLICANT: Tsuchihashi, Zenta

APPLICANT: Wolff, Roger K.

TITLE OF INVENTION: Megabase Transcript Map: No. 5872237e1

Sequences and Antibodies Thereeto

NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESSEE: TOWNSEND and TOWNSEND and CREW LLP
STREET: Two Embarcadero Center, 8th Floor
CITY: San Francisco
STATE: CA
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/724,394A
FILING DATE: 01-OCT-1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Fitts, Renee A.
REGISTRATION NUMBER: 35,136
REFERENCE/DOCKET NUMBER: 017957-000100
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-576-0200
TELEFAX: 415-576-0300
INFORMATION FOR SEQ ID NO: 21:
SEQUENCE CHARACTERISTICS:
LENGTH: 246240 base pairs
TYPE: nucleic acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..246240
OTHER INFORMATION: /note="HLA-H.CONTIG"
US-08-724-394A-21

Query Match 72.4%; Score 15.2; DB 2; Length 246240;
Best Local Similarity 85.0%; Pred. No. 86;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 aagtgactccctgactact 20
Db 139464 ATGTGACTGCTGCAACCT 139445

RESULT 10
US-08-724-394A-22/C
Sequence 22, Application US/08724394A
Patent No. 3672237
GENERAL INFORMATION:
APPLICANT: Feder, John N.
APPLICANT: Kronmal, Gregory S.
APPLICANT: Lauer, Peter M.
APPLICANT: Ruddy, David A.
APPLICANT: Thomas, Winston
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolf, Roger K.
TITLE OF INVENTION: Megabase Transcript Map: No. 5872237e1
TITLE OF INVENTION: Sequences and Antibodies Thereof
NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESSEE: TOWNSEND and TOWNSEND and CREW LLP
STREET: Two Embarcadero Center, 8th Floor
CITY: San Francisco
STATE: CA
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/724,394A
FILING DATE: 01-OCT-1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Fitts, Renee A.
REGISTRATION NUMBER: 35,136
REFERENCE/DOCKET NUMBER: 017957-000100
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-576-0200
TELEFAX: 415-576-0300
INFORMATION FOR SEQ ID NO: 22:
SEQUENCE CHARACTERISTICS:
LENGTH: 246240 base pairs
TYPE: nucleic acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..246240
OTHER INFORMATION: /note="HLA-H.CONTIG"
US-08-724-394A-22

Query Match 72.4%; Score 15.2; DB 2; Length 246240;
Best Local Similarity 85.0%; Pred. No. 86;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 aagtgactccctgactact 20
Db 139464 ATGTGACTGCTGCAACCT 139445

RESULT 11
US-08-417-492-1
Sequence 1, Application US/08417492
Patent No. 5750872
GENERAL INFORMATION:
APPLICANT: Bennett, Alan B.
APPLICANT: Brunnett, David A.
APPLICANT: Grantz, Alexander A.
TITLE OF INVENTION: Nucleic Acids Encoding Ascorbate Free
TITLE OF INVENTION: Radical Reductase and Their Uses
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew
STREET: One Market Plaza, Stewart Street Tower
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105-1492
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/417,492
FILING DATE: 05-APR-1995
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: Bastian, Kevin L.
REGISTRATION NUMBER: 34,774
REFERENCE/DOCKET NUMBER: 2307E-586US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-543-9600
TELEFAX: 415-543-5043
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1723 base pairs
TYPE: nucleic acid

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STRANDEDNESS: single
TOPOLOGY: linear
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: exon
LOCATION: 49..643
OTHER INFORMATION: /number- 1
FEATURE:
NAME/KEY: intron
LOCATION: 644..780
OTHER INFORMATION: /number- 1
FEATURE:
NAME/KEY: exon
LOCATION: 781..1484
OTHER INFORMATION: /number- 2
FEATURE:
NAME/KEY: CDS
LOCATION: join(49..643, 781..1487)
US-08-417-492-1

Query Match
Best Local Similarity 70.5%; Score 14.8; DB 1; Length 1723;
Best Local Similarity 88.9%; Pred. No. 68;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 tgaactccctgatacctt 21
DB 1424 TCACCCCTCTCTACTT 1441

RESULT 12
US-09-001-826-4
; Sequence 4, Application US/09001826A
; Patent No. 5965727
; GENERAL INFORMATION:
; APPLICANT: SONG, HEE-SOOK
; APPLICANT: BROTHERTON, JEFFREY E.
; APPLICANT: MIDHOLM, JACK M.
; TITLE OF INVENTION: SELECTABLE MARKER AND PROMOTER FOR PLANT TISSUE CULTURE
; FILE REFERENCE: 07001.C1
; CURRENT APPLICATION NUMBER: US/09/001,826A
; CURRENT FILING DATE: 1997-12-31
; EARLIER APPLICATION NUMBER: 08/937,739; 60/025,140
; EARLIER FILING DATE: 1997-07-25; 1996-07-26
; NUMBER OF SEQ ID NOS: 31
; SOFTWARE: Macintosh Wordperfect converted to PC ASCII Text
; SEQ ID NO: 4
; LENGTH: 2161
; TYPE: DNA (cDNA)
; ORGANISM: Nicotiana tabacum
US-09-001-826-4

Query Match
Best Local Similarity 70.5%; Score 14.8; DB 2; Length 2161;
Best Local Similarity 88.9%; Pred. No. 70;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 tgaactccctgatacctt 21
DB 593 tgaactccctgatacctt 610

RESULT 13
US-09-001-826-25
; Sequence 25, Application US/09001826A
; Patent No. 5965727
; GENERAL INFORMATION:
; APPLICANT: SONG, HEE-SOOK
; APPLICANT: BROTHERTON, JEFFREY E.
; APPLICANT: MIDHOLM, JACK M.
; TITLE OF INVENTION: SELECTABLE MARKER AND PROMOTER FOR PLANT TISSUE CULTURE
; FILE REFERENCE: TRANSFORMATION
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FILE REFERENCE: 07001.C1
; CURRENT APPLICATION NUMBER: US/09/001,826A
; CURRENT FILING DATE: 1997-12-31
; EARLIER APPLICATION NUMBER: 08/937,739; 60/025,140
; EARLIER FILING DATE: 1997-07-25; 1996-07-26
; NUMBER OF SEQ ID NOS: 31
; SOFTWARE: Macintosh Wordperfect converted to PC ASCII Text
; SEQ ID NO: 25
; LENGTH: 2161
; TYPE: DNA (cDNA)
; ORGANISM: Nicotiana tabacum
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 90..1940
US-09-001-826-25

Query Match
Best Local Similarity 70.5%; Score 14.8; DB 2; Length 2161;
Best Local Similarity 88.9%; Pred. No. 70;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 tgaactccctgatacctt 21
DB 593 tgaactccctgatacctt 610

RESULT 14
US-08-366-547-1
; Sequence 1, Application US/08366547
; Patent No. 5843737
; GENERAL INFORMATION:
; APPLICANT: Chen, Ian Bo
; APPLICANT: Bao, Shideng
; TITLE OF INVENTION: A NEW CANCER ASSOCIATED GENE, PROTEIN
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: DAVID G. CONLIN, DIKE, BRONSTEIN, ROBERTS
; ADDRESS: 6 CUSHMAN
; STREET: 130 Water Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/366,547
; FILING DATE: 30-DEC-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Eisenstein, Ronald I.
; REGISTRATION NUMBER: 30628
; REFERENCE/DOCKET NUMBER: 45072
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 523-3400
; TELEFAX: (617) 523-6440
; TELEX: 200291 STRE UR
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2652 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 120..2130
US-08-366-547-1
```


Query Match 70.5%; Score 14.8; DB 2; Length 2652;
 Best Local Similarity 88.9%; Pred. No. 72;

Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 agtgactcctgtctact 19
 |||||
 Db 2037 AGTGACTGCTGCTACT 2054

RESULT 15
 US-08-276-151-1/c

; Sequence 1, Application US/08276151
 ; Patent No. 5597719

; GENERAL INFORMATION:

; APPLICANT: Freed, Ellen

; APPLICANT: Ruggieri, Rosamaria

; TITLE OF INVENTION: Interaction of raf-1 and 14-3-3 proteins

; NUMBER OF SEQUENCES: 9

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Cooley Godward et al.

; STREET: Five Palo Alto Square

; CITY: Palo Alto

; STATE: CA

; COUNTRY: USA

; ZIP: 94036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/276,151

; FILING DATE: 14-JUL-1994

; CLASSIFICATION: 530

; ATTORNEY/AGENT INFORMATION:

; NAME: Torchia, Ph.D., Timothy E

; REGISTRATION NUMBER: 36,700

; REFERENCE/DOCKET NUMBER: ONTX-005/0005

; TELEPHONE: (415) 843-5481

; TELEFAX: (415) 857-0663

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 2977 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA to mRNA

; HYPOTHEICAL: NO

; ANTI-SENSE: NO

; ORIGINAL SOURCE:

; ORGANISM: Homo sapiens

; FEATURE:

; NAME/KEY: CDS

; LOCATION: 130..2076

; US-08-276-151-1

Query Match 70.5%; Score 14.8; DB 1; Length 2977;
 Best Local Similarity 88.9%; Pred. No. 74;

Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 tgaactcctgtactctt 21
 |||||
 Db 2436 TGAAGTCTGCTGCTACTT 2419

Search completed: July 25, 2001, 05:18:51
 Job time: 3214 sec

QY 3 tgcctcgtccagaggtt 19
|||||

Db 506 tgcctcgtccagaggtt 490

RESULT 5

LOCUS

AO068059

356 bp DNA

GSS

04-AUG-1998

DEFINITION

ACCSSION

AO068059

HE_2200_B2_F03_MF CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate-2200 Col-6 Row-L, DNA sequence.

VERSION

AO068059.1

GI:3379327

KEYWORDS

GSS.

SOURCE

human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 356)

AUTHORS

Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome

JOURNAL

MEDLINE

99380589

Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3887
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 2200 row: L column: 6
Class: BAC ends
High quality sequence stop: 356.

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

92 t

ORIGIN

Location/Qualifiers

1..356

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate-2200 Col-6 Row-L"

/clone_1lb="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coll DH10B"

BASE COUNT

111 a

86 c

67 g

Email: joan@udel.edu
Seq primer: T7.

FEATURES

source

Location/Qualifiers
1. 574
/organism="Gallus gallus"
/db_xref="taxon:9031"
/clone_pat="PK0004.12"
/clone_lib="chicken activated T cell cDNA"
/sex="male"
/cell_type="Con A-activated splenic T cell"
/lab_host="E. coli TOP10 F"
/note="Vector: pCDNA3"

BASE COUNT 165 a 130 c 154 g 125 t
ORIGIN

Query Match 86.3%; Score 16.4; DB 104; Length 574;
Best Local Similarity 94.4%; Pred. No. 6.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ttgtctctgcacagagt 18
|||||
Db 340 TTGCTCCTGCACAGCT 323

RESULT 8

A2664475 616 bp DNA GSS 14-DEC-2000
LOCUS IM0544H08R Mouse 10kb plasmid UUGCM library Mus musculus genomic
DEFINITION clone UUGCM0544H08 R, DNA sequence.
ACCESSION A2664475
VERSION A2664475.1 GI:11801621
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 616)
AUTHORS Dunn, D., Moyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A. and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0544 row: H column: 08
Seq primer: CACACAGAACACCTACAC
Class: plasmid ends
High quality sequence stop: 616.

FEATURES

source

Location/Qualifiers
1. 616
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGCM0544H08"
/clone_lib="Mouse 10kb plasmid UUGCM library"
/sex="Male"
/lab_host="E. coli strain XL10-gold, T1-resistant, F-"
/note="Vector: pMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA

BASE COUNT 156 a 162 c 131 g 167 t
ORIGIN

Query Match 86.3%; Score 16.4; DB 247; Length 616;
Best Local Similarity 94.4%; Pred. No. 6.8e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ttgtctctgcacagagt 19
|||||
Db 319 TTGCTCCTGCACAGCTT 336

RESULT 9

BE901069 680 bp mRNA EST 29-SEP-2000
LOCUS 601674415P1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3957315 5',
DEFINITION mRNA sequence.
ACCESSION BE901069
VERSION BE901069.1 GI:10389877
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 680)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-f@mail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: Image.llnl.gov
Plate: LCN834 row: n column: 04
High quality sequence start: 2
High quality sequence stop: 608.

FEATURES

source

Location/Qualifiers
1. 680
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3957315"
/clone_lib="NIH_MGC_21"
/tissue="choriocarcinoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: placenta; Vector: pOTB; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCCACG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 117 a 190 c 214 g 159 t
ORIGIN

Query Match 86.3%; Score 16.4; DB 141; Length 680;
 Best Local Similarity 94.4%; Pred. No. 6.9e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ttgtctcgcgcagaggt 18
 Db 371 TTGTCTCTCTCCAGAGCT 388

RESULT 10
 BG109825/c 978 bp mRNA EST 30-JAN-2001
 LOCUS 60280983F1 NIH_MGC-86 Homo sapiens cDNA clone IMAGE:4368405.5,
 DEFINITION mRNA sequence.
 ACCESSION BG109825
 VERSION BG109825.1 GI:1260331
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 REFERENCE 1 (bases 1 to 978)
 AUTHORS NIH-MGC <http://mgi.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LAM10022 row: f column: 22
 High quality sequence stop: 751.
 Location/Qualifiers
 1. 978
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_image="4368405"
 /clone_lib="NIH_MGC-86"
 /tissue_type="osteosarcoma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /note="Organ: bone; Vector: PCW-SPORT6, Site: 1; NotI;
 Site: 2; SalI; Cloned unidirectionally; oligo-dT primed.
 Average insert size 1.533 kb. Library enriched for
 full-length clones and constructed by Life Technologies.
 Note: this is a NIH-MGC Library."

BASE COUNT 250 a 250 c 276 g 202 t
 ORIGIN

Query Match 86.3%; Score 16.4; DB 173; Length 978;
 Best Local Similarity 94.4%; Pred. No. 7.3e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ttgtctcgcgcagaggt 18
 Db 844 TTGTCTCTCCAGAGCT 827

RESULT 11
 AI203923 206 bp mRNA EST 28-OCT-1998
 LOCUS qd7h07.x1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:1755069
 DEFINITION 3', mRNA sequence.
 ACCESSION AI203923
 VERSION AI203923.1 GI:3756529
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (bases 1 to 206)
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo
 , Ph.D.

FEATURES
 source
 1. 206
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_image="1755069"
 /clone_lib="Soares_testis_NHT"
 /sex="male"
 /lab_host="DH10B"
 /note="Vector: pT73D-Pac (Pharmacia) with a modified
 polylinker; Site: 1; Not I; Site: 2; Eco RI; 1st strand cDNA
 was prepared from mRNA obtained from Clontech Laboratories
 , Inc., and primed with a Not I - oligo(dT) primer [5'
 TGTTCACATCTGATGACGACGCGCCCAATTTTCTTTTCTTTT 3'].
 Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT73 vector. Library
 went through one round of normalization to C0t5, and was
 constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 50 a 48 c 53 g 55 t
 ORIGIN

Query Match 84.2%; Score 16; DB 17; Length 206;
 Best Local Similarity 100.0%; Pred. No. 9.1e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 4 gtcctcgcgcagaggt 19
 Db 52 GCTCTCTCCAGAGCT 67

RESULT 12
 AA504455 338 bp mRNA EST 18-AUG-1997
 LOCUS aa60a05.s1 NCI-CGAP_GCB1 Homo sapiens cDNA clone IMAGE:825296.3,
 DEFINITION mRNA sequence.
 ACCESSION AA504455
 VERSION AA504455.1 GI:2240615
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 338)
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
 Ph.D., Gerald Marti, M.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 320.
 Location/Qualifiers

FEATURES

source
 1. 338
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:825296"
 /clone_1lb="NCI-CGAP-C081"
 /tissue_type="colon tumor RER"
 /lab_host="DH10B"
 /note="Vector: p773D-Pac (Pharmacia) with a modified
 polylinker. Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
 was prepared from human tonsillar cells enriched for
 germinal center B cells by flow sorting (CD20⁺, IgD⁺),
 provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
 (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was
 primed with a Not I - oligo(dT) primer
 [5'-TGTTACCAATCTGAGTGGAGCGCGCCCTCTTTTCTTTTCTTTT-3',
 1. Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified p773 vector. Library
 went through one round of normalization, and was
 constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 79 a 85 c 82 g 95 t
 ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 341;
 Best Local Similarity 100.0%; Pred. No. 9.9e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 gctcctgcagaggtt 19
 |||||
 DB 86 gctcctgcagaggtt 101

RESULT 13
 AA577198 341 bp MRNA EST 12-SEP-1997
 LOCUS pm87b11.s1 NCI-CGAP-C09 Homo sapiens cDNA clone IMAGE:1075197 3'
 DEFINITION MRNA sequence.
 AA577198
 VERSION AA577198.1 GI:2354672
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 341)
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Ilan Kirsch, M.D., Michael R. Emmert-Buck, M.D.,
 Ph.D.

CDNA Library Preparation: M. Bento Soares, Ph.D.
 CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
 Insert length: 919 Std Error: 0.00
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 181.
 Location/Qualifiers

FEATURES

source
 1. 341
 /organism="Homo sapiens"

/db_xref="taxon:9606"
 /clone="IMAGE:1075197"
 /clone_1lb="NCI-CGAP-C09"
 /tissue_type="colon tumor RER"
 /lab_host="DH10B"
 /note="Organ: colon; Vector: p773D-Pac (Pharmacia) with a
 modified polylinker; 1st strand cDNA was prepared from
 RER+ colon tumor, and was then primed with a Not I -
 oligo(dT) primer. Double-stranded cDNA was ligated to Eco
 RI adaptors (Pharmacia), digested with Not I and cloned
 into the Not I and Eco RI sites of the modified p773
 vector. Library is not normalized. Library was
 constructed by Bento Soares and M. Fatima Bonaldo (Soares4
)."

BASE COUNT 79 a 85 c 82 g 95 t
 ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 341;
 Best Local Similarity 100.0%; Pred. No. 9.9e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 gctcctgcagaggtt 19
 |||||
 DB 189 gctcctgcagaggtt 204

RESULT 14
 A0751483 738 bp DNA GSS 19-JUL-1999
 LOCUS HS-5576_p2-B02-SPE RPCR-11 Human Male BAC Library Homo sapiens
 DEFINITION genomic clone Place-1152 Col-4 Row-D, DNA sequence.
 A0751483
 VERSION A0751483.1 GI:5538641
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 738)
 AUTHORS Mahairas G.G., Wallace J.C., Smith K., Swartzell S., Holzman T.,
 Mahairas G.G., Wallace J.C., Smith K., Swartzell S., Holzman T.,
 Keller A., Shaker R., Furlong J., Young J., Zhao S., Adams K.D. and
 Hood L.

Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome
 Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
 99380589
 COMMENT Contact: Mahairas G.G., Wallace J.C., Hood L.
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3687

Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCR-11. For BAC
 library availability, please contact Pieter de Jong
 (pieterdejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
 or from Research Genetics (<http://inforesgen.com>). BAC end Web Server:
<http://www.htsc.washington.edu>
 Plate: 1152 Row: D Column: 4
 Seq primer: SP6
 Class: BAC ends
 High quality sequence stop: 738.
 Location/Qualifiers

FEATURES

source
 1. 738
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Place-1152 Col-4 Row-D"
 /clone_1lb="RPCR-11 Human Male BAC Library"
 /sex="male"
 /note="Vector: pBAC3.6; Site 1: EcoRI; Site 2: EcoRI;
 Male blood DNA was isolated from one randomly chosen donor

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:23:04 ; Search time 247.87 Seconds

(without alignments)
50.664 Million cell updates/sec

Title: US-09-142-095-3

Sequence: 1 gcacagtgacacagtcacac 20

Scoring table:

IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Total number of hits satisfying chosen parameters: 1460202

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

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N_Geneseq_0601:*

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2: /SIDS1/gcgcdata/geneseq/NA1981.DAT:*

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20: /SIDS1/gcgcdata/geneseq/NA1999.DAT:*

21: /SIDS1/gcgcdata/geneseq/NA2000.DAT:*

22: /SIDS1/gcgcdata/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	AA79543	UGT1*1 gene exon 1
2	20	100.0	620	AA79540	Upstream DNA seqe
3	16.4	82.0	22	AA713057	3'-flanking region
4	15.8	79.0	27	AA576627	Collagen IV alpha
5	15.8	79.0	27	AA48039	Human Alport syndr
6	15.8	79.0	300	AA598515	Human cancer cell
7	15.8	79.0	392	AA50369	Human Goodpasture
8	15.8	79.0	507	AA50370	Human Goodpasture
9	15.8	79.0	680	AA50368	Human Goodpasture
10	15.8	79.0	685	AA50367	Human Goodpasture
11	15.8	79.0	738	AA57158	Human type IV coll

12	15.8	79.0	900	AA220991	Type IV collagen N
13	15.8	79.0	900	AA80993	Human alpha3(IV)NC
14	15.2	76.0	1276	AA713921	Aspergillus oryzae
15	15.2	76.0	1801	AA635997	Arabidopsis thalia
16	15.2	76.0	1803	AA631372	Arabidopsis thalia
17	15	75.0	2530	AA638424	Arabidopsis thalia
18	14.8	74.0	393	AA666883	Novel human polynu
19	14.8	74.0	430	AA719914	Human gene signatu
20	14.8	74.0	519	AA696028	Human breast tumor
21	14.8	74.0	519	AA696028	DNA molecule encod
22	14.8	74.0	887	AA60894	Sequence of plasmid
23	14.8	74.0	1100	AA60127	Sequence encoding
24	14.8	74.0	1100	AA60103	Sequence of human
25	14.8	74.0	1101	AA603003	Sequence including
26	14.8	74.0	1101	AA60103	Sequence of human
27	14.8	74.0	1101	AA60103	Human immunointerf
28	14.8	74.0	1144	AA60240	Sequence encoding
29	14.8	74.0	1156	AA60240	Human recombinant
30	14.8	74.0	1161	AA60327	Human interferon-g
31	14.8	74.0	1161	AA60103	Sequence of the pl
32	14.8	74.0	1210	AA60055	Human interferon-g
33	14.8	74.0	1226	AA604819	Sequence of the IF
34	14.8	74.0	1350	AA60335	Sequence of plasmid
35	14.8	74.0	2059	AA60895	Nucleotide sequenc
36	14.8	74.0	2171	AA60895	Nucleotide sequenc
37	14.8	74.0	2274	AA60895	Protoposin cDNA.
38	14.8	74.0	2740	AA60895	Human prosaposin c
39	14.8	74.0	2749	AA60893	Sequence of plasmid
40	14.8	74.0	2827	AA60893	Sequence of plasmid
41	14.8	74.0	3000	AA60892	Arabidopsis thalia
42	14.8	74.0	5274	AA642949	Arabidopsis thalia
43	14.8	74.0	5319	AA65069	Human interferon-g
44	14.8	74.0	5961	AA65069	Human chromosome 6
45	14.8	74.0	68940	AA57351	

ALIGNMENTS

AA79543	1	UGT1*1 gene exon 1 upstream PCR primer C.
AA79543	23-JAN-1998	(first entry)
AA79543	23-JAN-1998	Uridine diphosphate glucuronosyltransferase gene; UGT;
AA79543	23-JAN-1998	Gilbert's syndrome; GS; unconjugated hyperbilirubinemia;
AA79543	23-JAN-1998	bilirubin glucuronidation; Crigler-Najjar; type 2; drug metabolism;
AA79543	23-JAN-1998	Drug trial efficiency; screening; PCR primer; ss.
AA79543	23-JAN-1998	Synthetic.
AA79543	23-JAN-1998	Homo sapiens.
AA79543	23-JAN-1998	W09732042-A2.
AA79543	23-JAN-1998	04-SEP-1997.
AA79543	23-JAN-1998	03-MAR-1997.
AA79543	23-JAN-1998	97MO-GB00577.
AA79543	23-JAN-1998	16-MAR-1996.
AA79543	23-JAN-1998	96GB-0005598.
AA79543	23-JAN-1998	01-MAR-1996.
AA79543	23-JAN-1998	96GB-0004480.
AA79543	23-JAN-1998	(UTD-) UNIV DUNDEE.
AA79543	23-JAN-1998	Burchell B.
AA79543	23-JAN-1998	WPI: 1997-448702/41.
AA79543	23-JAN-1998	Improving drug trial efficiency comprises identifying participants

PT with Gilbert's syndrome - useful as their altered drug metabolism
 PT may hinder result interpretation
 PS Claim 14; Page 12; 31pp; English.

CC This PCR primer (with primers AAT79542-44) flanks the TATA box sequence
 CC upstream of the uridine diphosphate glucuronosyltransferase (UGT) gene
 CC 141 exon 1 (see AAT79540), and was used to amplify fragments of 98-100
 CC bp. This gene is known to be associated with Gilbert's syndrome (GS). GS
 CC is a mild, common form of unconjugated hyperbilirubinaemia associated
 CC with reduced bilirubin glucuronidation capacity. Analysis of the genetic
 CC basis of GS has allowed 2 forms to be identified. One is a mild form
 CC associated with a homozygous 2 bp insertion in the TATA sequence
 CC upstream of the UGT*1 exon 1, and the other is a more severe form
 CC associated with heterozygosity for a mutation which, when homozygous,
 CC causes Crigler-Najjar type 2 disease. The first form is autosomal
 CC recessive and the second is inherited dominantly. Patients suffering from
 CC GS, which is benign, may have altered metabolism of some drugs, making it
 CC difficult to determine if an effect is due to the drug or the syndrome.
 CC Drug trial efficiency would be improved if potential participants can be
 CC screened for the genetic basis of GS, and eliminated or included on
 CC basis of them possessing or not possessing GS.

XX Sequence 20 BP; 7 A; 6 C; 4 G; 3 T; 0 other;

Query Match 100.0%; Score 20; DB 18; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.11; 0; Indels 0; Gaps 0;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gtcacgtgacacagtcacac 20
 |||||
 Db 1 gtcacgtgacacagtcacac 20

RESULT 2

AAT79540
 ID AAT79540 standard; DNA; 620 BP.

XX AC AAT79540;

XX DT 23-JAN-1998 (first entry)

XX DE Upstream DNA sequence of UGT1*1 gene exon 1.

XX KW Uridine diphosphate glucuronosyltransferase gene; UGT;

KW Gilbert's syndrome; GS; unconjugated hyperbilirubinaemia;
 KM Bilirubin glucuronidation; Crigler-Najjar; type 2; drug metabolism;

KM Drug trial efficiency; screening; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers
 FT protein_bind 2..8
 FT /tag- a
 FT /bound_moelty- APl

FT GC_signal 93..102
 FT /tag- b
 FT /note- "SPL binding site"

FT misc-feature 180..191
 FT /tag- c
 FT /note- "feature indicated in patent, but no further
 explanation is given"

FT protein_bind 316..323
 FT /tag- d
 FT /bound_moelty- AP3

FT protein_bind 317..324
 FT /tag- e
 FT /bound_moelty- CLBP

FT misc-feature 366..376
 FT /tag- f
 FT /note- "feature indicated in patent, but no further
 explanation is given"

FT misc-feature 366..376
 FT /tag- f
 FT /note- "feature indicated in patent, but no further
 explanation is given"

FT protein_bind 386..392

FT /tag- g

FT /bound_moelty- APl

FT protein_bind 513..519

FT /tag- h

FT /bound_moelty- APl

FT misc-feature 513..520

FT /tag- i

FT /note- "feature indicated in patent, but no further
 explanation is given"

FT protein_bind 520..532

FT /tag- j

FT /bound_moelty- HNF1

FT TATA_signal 558..572

FT /tag- k

FT /note- "corresponds to positions -53 to -39 in patent"

FT misc-feature 595

FT /tag- l

FT /note- "feature indicated in patent, but no further
 explanation is given"

FT CDS 612

FT /tag- m

FT /codon_start- 612

XX WO9732042-A2.

XX PD 04-SEP-1997.

XX PF 03-MAR-1997; 97WO-GB00577.

XX PR 16-MAR-1996; 96GB-0005598.

XX PR 01-MAR-1996; 96GB-0004480.

XX PA (UYDU-) UNIV DUNDEE.

XX PI Burchell B;

XX DR WPI, 1997-448702/41.

XX PT Improving drug trial efficiency comprises identifying participants
 with Gilbert's syndrome - useful as their altered drug metabolism
 may hinder result interpretation

XX PS Claim 11; Fig 4; 31pp; English.

XX CC This sequence represents the upstream sequence, positions -61 to 9
 in the patent, of uridine diphosphate glucuronosyltransferase (UGT)
 gene 1*1 exon 1. This gene is known to be associated with Gilbert's
 syndrome (GS). GS is a mild, common form of unconjugated
 hyperbilirubinaemia associated with reduced bilirubin glucuronidation
 capacity. Analysis of the genetic basis of GS has allowed 2 forms to be
 identified. One is a mild form associated with a homozygous 2 bp
 insertion in the TATA sequence upstream of the UGT1*1 exon 1, and
 the other is a more severe form associated with heterozygosity for a
 mutation which, when homozygous, causes Crigler-Najjar type 2 disease.
 The first form is autosomal recessive and the second is inherited
 dominantly. Patients suffering from GS, which is benign, may have
 altered metabolism of some drugs, making it difficult to determine if an
 effect is due to the drug or the syndrome. Drug trial efficiency would
 be improved if potential participants can be screened for the genetic
 basis of GS, and eliminated or included on basis of them possessing or
 not possessing GS. In this case, screening involves PCR amplification
 of the UGT gene, using the primers described in AAT79541-44.

XX Sequence 620 BP; 157 A; 127 C; 151 G; 185 T; 0 other;

Query Match 100.0%; Score 20; DB 18; Length 620;
 Best Local Similarity 100.0%; Pred. No. 0.19;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gtcacgtgacacagtcacac 20
 |||||

Db 508 gtcacgtgacacagtcacaac 527

RESULT 3

AAV13057/c
ID AAV13057 standard; DNA; 22 BP.

XX AAV13057;

XX 18-MAY-1998 (first entry)

XX 3'-flanking region of an interleukin 6 gene PCR primer.

XX Interleukin 6; IL-6; PCR primer; genotype; osteoporosis; human;

XX polymorphism; ss.

XX Synthetic.

XX Homo sapiens.

XX W09743446-A2.

XX 20-NOV-1997.

XX 16-MAY-1997; 97WO-GB01337.

XX 16-MAY-1996; 96GB-0010281.

XX (GEMI-) GEMINI INT HOLDINGS LTD.

XX Grant SFA, Raiston SH;

XX WPI; 1998-008905/01.

XX Determination of genotype of an IL-6 gene in an individual - used

XX for diagnosis of or identification of a predisposition or

XX susceptibility to a disease, particularly osteoporosis

XX Claim 9; Page 6; 20pp; English.

XX The present sequence represents a PCR primer used in the present

XX invention. The present invention describes an in vitro method of

XX diagnosis of a disease in an animal. The method comprises determining

XX the genotype of an interleukin-6 (IL-6) gene in the animal. The

XX invention also describes: (1) a method of identifying an animal

XX predisposed or susceptible to a disease, comprising determining the

XX genotype of an IL-6 gene in the animal; (2) a composition for use in

XX diagnosing a disease in an animal, the composition comprising one or

XX more primer nucleic acid molecules adapted to amplify a portion of a

XX 3' flanking region of an IL-6 gene in the animal; and (3) a composition

XX for use in identifying an animal predisposed or susceptible to a

XX disease, the composition comprising one or more primer nucleic acid

XX molecules adapted to amplify a portion of a 3' flanking region of an

XX IL-6 gene in the animal. The method can be used for the diagnosis of

XX or identifying predisposition or susceptibility to a disease,

XX particularly osteoporosis.

XX Sequence 22 BP; 4 A; 4 C; 7 G; 7 T; 0 other;

Query Match 82.0%; Score 16.4; DB 19; Length 22;

Best Local Similarity 94.4%; Pred. No. 9;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaac 19

Db 19 TCACGTGACACAGTCACAA 2

RESULT 4

AAZ57627
ID AAZ57627 standard; DNA; 27 BP.

XX AAZ57627;

XX 05-APR-2000 (first entry)

XX Collagen IV alpha 3 deletion mutant fragment #1.

XX Collagen IV alpha 3; deletion mutant; nucleotide probe; genomic profile;

XX Alport syndrome; disease prognosis; health screening;

XX health management tool; ss.

XX Unidentified.

XX W0964626-A2.

XX 16-DEC-1999.

XX 04-JUN-1999; 99WO-GB01779.

XX 06-JUN-1998; 96GB-0012098.

XX 23-DEC-1998; 98GB-0028289.

XX (GENO-) GENOSTIC PHARMA LTD.

XX Roberts GW;

XX WPI; 2000-097546/08.

XX Nucleotide probes used in genetic screens for determining genomic

XX profiles, e.g. for prognosis or management -

XX Example 5; Page 25; 14pp; English.

XX This sequence is an collagen IV alpha 3 deletion mutant fragment. The

XX mutated gene is an example of a variant that can be detected using the

XX new nucleotide probes of the invention. The probes consist of

XX complementary DNA or RNA for the detection of variants in a group of

XX target genes associated with particular diseases. The mutant variant

XX represented by this sequence gives rise to Alport syndrome phenotype. The

XX specification lists approximately 2500 genes that are target groups for

XX the nucleotide probes of the invention. The probes are used to determine

XX the genomic profile of an individual, particularly for:

XX (a) prognosis and management of disease (or determining susceptibility

XX to disease);

XX (b) predicting a patient's response to therapy and symptom profiles;

XX (c) for health screening;

XX (d) to develop new strategies for therapy and clinical trials; and

XX (e) to construct health management algorithms or models, e.g. of the

XX likelihood of developing disease.

XX Antibodies specific for the proteins encoded by the genes in the target

XX list, can be used similarly, identification of a core group of genes

XX associated with disease, makes possible the use of genetic profiling as a

XX universal health management tool.

XX Sequence 27 BP; 7 A; 10 C; 3 G; 7 T; 0 other;

Query Match 79.0%; Score 15.8; DB 21; Length 27;

Best Local Similarity 89.5%; Pred. No. 19;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaac 20

Db 8 TCACGTGACACAGTCACAAC 26

RESULT 5

AAZ48039
ID AAZ48039 standard; DNA; 27 BP.

XX AAZ48039;

XX 08-MAR-2000 (first entry)

XX Human Alport syndrome oligonucleotide #1.

XX Human; glycogen storage disease 2; genetic filling; identification;
 KW polymorphic variation; mutation; detection; probe; prognosis; headache;
 KW gene association; health screening; therapy; clinical trial;
 KW absorption; distribution; metabolism; elimination; oncology; dementia;
 KW central nervous system; behaviour; brain injury; psychosis; personality;
 KW cardiovascular; gastrointestinal; respiratory; immunology; development;
 KW skin; bone; muscle; endocrine; sexual dysfunction; ss.
 XX Homo sapiens.
 OS
 PN MO9964627-A2.
 XX
 PD 16-DEC-1999.
 XX
 PF 04-JUN-1999; 99WO-GB01780.
 XX
 PR 06-JUN-1998; 98GB-0012099.
 PR 20-JUN-1998; 98GB-0013291.
 PR 24-JUN-1998; 98GB-0013611.
 PR 27-JUN-1998; 98GB-0013835.
 PR 01-JUL-1998; 98GB-0014110.
 PR 07-JUL-1998; 98GB-0014580.
 PR 16-JUL-1998; 98GB-0015438.
 PR 18-JUL-1998; 98GB-0015574.
 PR 24-JUL-1998; 98GB-0015576.
 PR 24-JUL-1998; 98GB-0016085.
 PR 05-AUG-1998; 98GB-0016921.
 PR 07-AUG-1998; 98GB-0017097.
 PR 08-AUG-1998; 98GB-0017200.
 PR 14-AUG-1998; 98GB-0017632.
 PR 19-AUG-1998; 98GB-0017943.
 XX
 XX (GENO-) GENOSTIC PHARMA LTD.
 XX
 PI Roberts GM;
 DR MPI: 2000-097547/08.
 XX
 PT Nucleotide probes used in genetic screens for determining genomic
 PT profiles, e.g. for prognosis or management -
 PS Example 5; Page 25; 745pp; English.
 XX
 CC The present invention describes new nucleotide probes (i) comprising
 CC complementary DNA or RNA, used in the detection of variants in a group
 CC of target genes associated with particular diseases. Detected variants
 CC are mutations or polymorphisms. Included in the specification are lists
 CC of genes for the following target groups: (i) ADME (absorption,
 CC distribution, metabolism and elimination - about 1000 genes);
 CC (ii) oncology (about 1200 genes); (iii) central nervous system (about
 CC 850 genes); (iv) dementia (about 250 genes); (v) brain injury (about
 CC personality (about 500 genes); (vii) cardiovascular (about 1100 genes);
 CC (ix) gastrointestinal (about 1100 genes); (x) respiratory (about 800
 CC genes); (xi) immunology (about 1100 genes); (xii) development (about 2000
 CC and endocrine (about 1100 genes); (xv) headache (about 250 genes); and
 CC genomic profile of an individual, particularly for: (a) prognosis and
 CC management of disease (or determining susceptibility to disease);
 CC (b) predicting a patient's response to therapy and symptom profiles;
 CC (c) for health screening; (d) to develop new strategies for therapy and
 CC clinical trials; and (e) to construct health management algorithms or
 CC models, e.g. of the likelihood of developing disease. The present
 CC sequence represents an oligonucleotide used to illustrate a deletion in
 CC Alport syndrome, which is given in an example from the present invention.
 XX
 SQ Sequence 27 BP; 7 A; 10 C; 3 G; 7 T; 0 other;

Query Match

79.0%; Score 15.8; DB 21; Length 27;

Best Local Similarity 89.5%; Pred. No. 19;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 tcacgtgcacacgtcaaac 20
 Db ||||| ||||| |||||
 8 tcacgcgcacacgtcaaac 26

RESULT 6
 AAX98515/C
 ID AAX98515 standard; cDNA: 300 BP.
 AC AAX98515;
 XX
 DT 24-SEP-1999 (first entry)
 XX
 DE Human cancer cell derived cDNA #241.
 XX
 KW Cancer; human; colon; breast; lung; transmembrane receptor; ATPase;
 KW integral membrane protein; aspartyl protease; GATA family; wt family;
 KW transcription factor; G-protein alpha subunit; protein phosphatase;
 KW photoester binding protein; diacylglycerol binding protein; trypsin;
 KW protein kinase; tyrosine phosphatase; developmental signalling protein;
 KW Wt/rsp5/WMP domain; therapy; forensic; genetic mapping; diagnostic;
 KW Wilm's tumour; retinoblastoma; sarcoma; colorectal adenocarcinoma;
 KW leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal;
 KW prostate; ss.
 KW
 KW Homo sapiens.
 OS
 XX
 PN WO9933982-A2.
 XX
 PD 08-JUL-1999.
 XX
 PF 22-DEC-1998; 98WO-US27610.
 XX
 PR 21-DEC-1998; 98US-0217471.
 PR 23-DEC-1997; 97US-0068755.
 PR 03-APR-1998; 98US-0080664.
 PR 21-OCT-1998; 98US-0105234.
 PR 27-OCT-1998; 98US-0105877.
 XX
 PA (CHIR) CHIRON CORP.
 PI (HYSE-) HXSEQ INC.
 XX
 PI Crivenjakov R, Dickson M, Drmanac R, Drmanac S;
 PI Escobedo J, Garcia PD, Garcia V, Ciese K, Innis MA;
 PI Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;
 PI Lanson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
 PI Stache-Crain B, Sudduth-Klinger J, Williams LT;
 PI MPI: 1999-430243/36.
 DR
 XX
 XX New isolated human polynucleotides
 PS Claim 1; Page 375; 591pp; English.
 XX
 CC This invention describes novel isolated human polynucleotides obtained
 CC by screening for differential expression in colon cancer, breast cancer
 CC and lung cancer cell lines. The polynucleotides of the invention are
 CC represented in AAX98275-X9918 and encode polypeptides of protein
 CC families selected from 4 transmembrane segments integral membrane
 CC proteins, 7 transmembrane receptors, ATPases associated with various
 CC cellular activities (AA), eukaryotic aspartyl proteases, GATA family of
 CC transcription factors (AA), tyrosine phosphatase, trypsin, wt family of
 CC diacylglycerol binding proteins, protein kinase, protein phosphatase 2C,
 CC signalling proteins and Wt/rsp5/WMP domain containing proteins. The
 CC encoded polypeptides also have a functional domain selected from Ank
 CC bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger
 CC (C2H2 type), zinc finger (C2HC class), and zinc-binding metalloprotease

```

Query Match      . 79.0%; Score 15.8; DB 20; Length 300;
Best Local Similarity 89.5%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0

```

RESULT	7
AAA50369	
ID	AAA50369 standard; DNA; 392 BP

DT 20-NOV-2000 (first entry)

DE Human Goodpasture antigen Deltaii/IV/V cDNA

KM Goodpasture antigen; GPdelta11/IV/V; human; GPP;
 KM goodpasture antigen binding protein; autoimmune disease; apoptosis
 KM cancer; tumour; gene therapy; ss.

OS Homo sapiens

Key	Location/Qualifiers
1	310

/*tag= a

人々

PN MOZ0005060/-AZ.
XX

FD-31-AUG-2000
XX

24 FEB 2000, 2000M0-1800024
11
17
XX

44 FEB 23 1955
XX XX

XX

XX 1 0 0 0 0

DR P-PSDB; AAY95920.

Novel Goodpasture and









CC antigen (GP) Delta:

Figure 1

50 sequence 392 BP; 134 A; 80 C; 81 G; 97 T; 0 other,

Query Match	79.0%;	Score 15.8;	DB 21;	Length 392;
Best Local Similarity	89.5%;	Pred. No. 29;		
Matches 17; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

RESULT	8
AAA50370	
ID	AAA50370 standard; DNA; 507 BP.

AC AAA50370;

DT 20-NOV-2000 (first entry)

Human Goodpasture antigen DeltailV cDNA

KM Goodpasture antigen; GPdeltaIII/V; human; GPPB; Goodpasture antigen binding protein; autoimmunity; disease; resistance

XY cancer; tumour; gene therapy; ss

hollo sapiens
xy

localtion/qualifiers
1-219

Met-deleted GPdeltaIII/V

⊗ =

[illegible]

XX 5
0
1
5
6
6
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4
4
2

XX

XX.

DR P-PSDB; AAY95921

PT Novel Goodpasture

XX

XX

antigen (GP) DeltaII/V (see AAY95921), an alternative form of

CDNA was obtained by subcloning a cDNA encoding the protein into a

22Y95900-11) which bind to and phosphorylate the unique N-terminal residues of liver glucoproteins (GPIs, see

cc region of Indian or, and much are highly expressed in several

CC autoimmune conditions. Claimed methods for treating an autoimmune
 CC disorder, cell apoptosis or a tumour involve modifying the
 CC expression or activity of GPBP, especially using a GP-derived
 CC peptide, such as GPdeltaII/V or a nucleic acid sequence encoding
 CC it.

XX Sequence 507 BP; 159 A; 113 C; 104 G; 131 T; 0 other;

Query Match 79.0%; Score 15.8; DB 21; Length 507;
 Best Local Similarity 89.5%; Pred. No. 31;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaac 20
 Db 65 tcacccgacacagtcacaac 83

RESULT 9

AAA50368 standard; DNA; 680 BP.

AC AAA50368;

DT 20-NOV-2000 (first entry)

DE Human Goodpasture antigen DeltaIII cDNA.

XX Goodpasture antigen; GPdeltaIII; human; GPBP;

KW goodpasture antigen binding protein; autoimmune disease; apoptosis;

XX cancer; tumour; gene therapy; ss.

OS Homo sapiens.

XX Key

FT CDS

FT Location/Qualifiers

FT 1..219

FT /*tag- a

FT /product- Met-deleted GPdeltaIII

FT /partial

XX MO200050607-A2.

XX 31-AUG-2000.

XX 24-FEB-2000; 2000MO-IB00324.

XX 24-FEB-1999; 99US-0121483.

XX (SAUS/) SAUS J.

XX Saus J;

XX WPI: 2000-572094/53.

XX P-PSDB: AAY95919.

XX Novel Goodpasture antigen binding proteins useful for diagnosing and

XX treating autoimmune disorders, tumor, and preventing cell apoptosis

XX Claim 25; Page 152-153; 158pp; English.

XX The present sequence is that of cDNA encoding human Goodpasture

XX antigen (GP) DeltaIII (see AAY95919), i.e. an alternative form of

XX human GP resulting from splicing out of exon III. The cDNA was

XX obtained by subcloning a cDNA encoding the protein into a modified

XX pET15b vector including an initiator Met. The invention relates

XX to novel Goodpasture antigen binding proteins (GPBPs, see

XX AAY95900-11), which bind to and phosphorylate the unique N-terminal

SO Sequence 680 BP; 204 A; 159 C; 145 G; 172 T; 0 other;

Query Match 79.0%; Score 15.8; DB 21; Length 680;
 Best Local Similarity 89.5%; Pred. No. 32;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaac 20
 Db 65 tcacccgacacagtcacaac 83

RESULT 10

AAA50367 standard; DNA; 685 BP.

AC AAA50367;

DT 20-NOV-2000 (first entry)

DE Human Goodpasture antigen DeltaV cDNA.

XX Goodpasture antigen; GPdeltaV; goodpasture antigen binding protein;

KW GPBP; human; autoimmune disease; apoptosis; cancer; tumour;

XX gene therapy; ss.

OS Homo sapiens.

XX Key

FT CDS

FT Location/Qualifiers

FT 1..636

FT /*tag- a

FT /product- Met-deleted GPdeltaV

FT /partial

XX MO200050607-A2.

XX 31-AUG-2000.

XX 24-FEB-2000; 2000MO-IB00324.

XX 24-FEB-1999; 99US-0121483.

XX (SAUS/) SAUS J.

XX Saus J;

XX WPI: 2000-572094/53.

XX P-PSDB: AAY95918.

XX Novel Goodpasture antigen binding proteins useful for diagnosing and

XX treating autoimmune disorders, tumor, and preventing cell apoptosis

XX Claim 25; Page 150-151; 158pp; English.

XX The present sequence is that of cDNA encoding human Goodpasture

XX antigen (GP) DeltaV (see AAY95918), i.e. an alternative form of

XX human GP resulting from splicing out of exon V. The cDNA was

XX obtained by subcloning a cDNA encoding the protein into a modified

XX pET15b vector including an initiator Met. The invention relates

XX to novel Goodpasture antigen binding proteins (GPBPs, see

XX AAY95900-11), which bind to and phosphorylate the unique N-terminal

XX autoimmune conditions. Claimed methods for treating an autoimmune

XX disorder, cell apoptosis or a tumour involve modifying the

XX expression or activity of GPBP, especially using a GP-derived

XX peptide, such as GPdeltaV or a nucleic acid sequence encoding it.

XX Sequence 685 BP; 206 A; 157 C; 138 G; 184 T; 0 other;

XX Query Match 79.0%; Score 15.8; DB 21; Length 685;
 Best Local Similarity 89.5%; Pred. No. 32;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

2 tcacgtgacacagtcacaac 20
||||| |||||||
Db 65 tcacccgacacagtcacaac 83

RESULT 11

AA257158
ID AA257158 standard; CDNA; 738 BP.

AA257158;

03-APR-2000 (first entry)

Human type IV collagen alpha 3 chain nucleotide sequence SEQ ID NO:9.

Human; type IV collagen; anti-angiogenic; angiogenesis; cancer;
benign tumour; rheumatoid arthritis; diabetic retinopathy; psoriasis;
myocardial angiogenesis disease; Osler-Webber Syndrome; telangiectasia;
atherosclerosis; scleroderma; hypertrophic scar; cat scratch disease;
contraception; obesity; ss.

Homo sapiens.

W09965940-A1.

23-DEC-1999.

17-JUN-1999; 99MO-US13737.

17-JUN-1998; 98US-0089689.

25-MAR-1999; 99US-0126175.

(BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

Kalluri R;

WPI: 2000-097708/08.

P-PSDB; AA167942.

Anti-angiogenic proteins comprising the NC1 domain of the alpha 1, 2 or 3 chain of type IV collagen used in, e.g. treatment of benign tumors and rheumatoid arthritis -

Claim 33; Fig 16A; 117pp; English.

The present sequence encodes the human type IV collagen alpha 3 chain. The present invention describes an isolated protein chosen from the NC1 domain of the alpha 1, alpha 2 or alpha 3 chains of type IV collagen or a fragment, analogue, derivative or mutant, which has anti-angiogenic properties. The anti-angiogenic proteins, multimers and chimeras are useful for inhibiting angiogenic activity in mammalian tissue, especially for treating diseases chosen from angiogenesis-dependent cancers, benign tumors, rheumatoid arthritis, diabetic retinopathy, psoriasis, ocular angiogenesis diseases, Osler-Webber Syndrome, myocardial angiogenesis, plaque neovascularisation, telangiectasia, haemophilic joints, angiodioma, wound granulation, intestinal adhesions, atherosclerosis, scleroderma, hypertrophic scars, cat scratch disease, Helicobacter pylori ulcers, dialysis graft vascular access stenosis, contraception and obesity. The compositions can be used to inhibit a disease characterised by angiogenic activity, in conjunction with radiation therapy, chemotherapy or immunotherapy.

Sequence 738 BP; 204 A; 184 C; 162 G; 188 T; 0 other;

Query Match 79.0%; Score 15.8; DB 21; Length 738;
Best Local Similarity 89.5%; Pred. No. 32;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Db 68 tcacccgacacagtcacaac 86

RESULT 12

AA220091
ID AA220091 standard; DNA; 900 BP.

AA220091;

05-JAN-2000 (first entry)

Type IV collagen NC1 domain alpha-3 monomer DNA.

Type IV collagen; NC1 domain; non-collagenous domain; human;
angiogenesis; tumour; metastasis; therapy; diabetic retinopathy;
rheumatoid arthritis; retinal neovascularization;
choroidal neovascularization; macular degeneration;
corneal neovascularization; retinopathy of prematurity;
corneal graft rejection; neovascular glaucoma;
retrolental fibroplasia; epidemic keratoconjunctivitis;
vitamin A deficiency; contact lens overwear; atopic keratitis;
superior limbic keratitis; pterygium keratitis sicca; sogrens;
acne rosacea; phlyctenulosis; syphilis; Mycobacteria infection;
lipid degeneration; chemical burn; herpes simplex infection;
Herpes zoster infection; protozoan infection; Kaposi's sarcoma;
Mooren ulcer; Terrien's marginal degeneration;
marginal keratolysis; trauma; systemic lupus; polyarteritis;
Wegener's sarcolidosis; scleritis; Steven's Johnson disease;
radial keratotomy; sickle cell anaemia; sarcoid;
pseudoxanthoma elasticum; Paget's disease; vein occlusion;
artery occlusion; carotid obstructive disease; chronic uveitis;
chronic vitritis; Lyme's disease; Eales disease; Beethers disease;
myopia; optic pty; Stargart's disease; pars planitis;
chronic retinal detachment; hypervascosity syndrome; toxoplasmosis;
post-laser complication; fibrovascular tissue proliferation;
haemangioma; Osler-Webber-Rendu; AIDS; ocular neovascular disease;
osteoarthritis; chronic inflammation; Crohn's disease;
ulcerative colitis; psoriasis; atherosclerosis; pemphigoid; ss.

Homo sapiens.

Synthetic.

Key Location/Qualifiers

FT CDS 40..846

FT sig_peptide 40..90

FT mat_peptide 91..843

FT /tag- b

FT /product- "BM40 signal peptide"

FT /tag- c

FT /product- "affinity-tagged alpha-3 monomer"

PN W09949885-A2.

XX 07-OCT-1999.

PD 26-MAR-1999; 99MO-US06445.

PR 27-MAR-1998; 98US-0079783.

PR 29-OCT-1998; 98US-0106170.

XX (UNITV) UNIV KANSAS MEDICAL CENT.

XX Hudson BG, Sarraf MP;

XX WPI, 1999-601287/51.

XX P-PSDB; AA131993.

XX Inhibition of angiogenesis with non-collagenous alpha chain monomer

XX useful for treating e.g. tumor growth or metastasis,

XX neovascularisation, etc -

XX Disclosure; Fig 17c/ 56pp; English.

XX This is the nucleotide sequence of a recombinant DNA encoding a
 CC type IV collagen non-collagenous (NC1) domain alpha-3 polypeptide
 CC (see AAY1393) composed of a BM40 signal sequence (which is cleaved
 CC from the mature protein) to facilitate protein secretion, and a
 CC mature protein comprising an affinity tag (facilitates purification
 CC and identification of the material) and the alpha-1 chain monomer.
 CC The invention provides methods and kits for inhibiting angiogenesis,
 CC tumour growth and metastasis, and endothelial cell interaction with
 CC the extracellular matrix, each method comprising contacting the
 CC tumour or animal tissue with 1 or more isolated type IV collagen
 CC NC1 alpha chain monomer(s) selected from the group consisting of
 CC alpha-1, alpha-2, alpha-3 and alpha-6 NC1 chain monomers. The
 CC monomers can be produced via recombinant protein expression. The
 CC polynucleotides and polypeptides are used to treat an angiogenesis-
 CC mediated disorder or condition, especially selected from solid and
 CC blood-borne tumours, diabetic retinopathy, rheumatoid arthritis,
 CC retinal neovascularization, choroidal neovascularization, macular
 CC degeneration, corneal neovascularization, retinopathy of prematurity,
 CC corneal graft rejection, neovascular glaucoma, retrolental
 CC fibroplasia, epidemic keratoconjunctivitis, vitamin A deficiency,
 CC contact lens overwear, atopic keratitis, superior limbic keratitis,
 CC pterygium keratitis sicca, seborrheic acne rosacea, phlyctenulosis,
 CC syphilis, mycobacteria infections, lipid degeneration, chemical
 CC burns, bacterial ulcers, fungal ulcers, herpes simplex infections,
 CC herpes zoster infections, protozoan infections, Kaposi's sarcoma,
 CC Mooren ulcer, Terrien's marginal degeneration, marginal keratolysis,
 CC leucoma, systemic lupus, polyarteritis, Wegener's granulomatosis,
 CC scleritis, Steven's Johnson disease, radial keratotomy, sickle cell
 CC anemia, sarcoid, pseudoxanthoma elasticum, Paget's disease, vein
 CC occlusion, artery occlusion, carotid obstructive disease, chronic
 CC uveitis, chronic vitritis, Lyme's disease, Balo's disease, Behcet's
 CC disease, myopia, optic pits, Stargardt's disease, pars planitis,
 CC chronic retinal detachment, hyperviscosity syndromes, toxoplasmosis,
 CC post-laser complications, abnormal proliferation of fibrovascular
 CC tissue, haemangiomas, Osler-Weber-Rendu, AIDS, ocular neovascular
 CC disease, osteoarthritis, chronic inflammation, Crohn's disease,
 CC ulcerative colitis, psoriasis, atherosclerosis, and pemphigoid (all
 CC claimed).

SQ Sequence 900 BP; 228 A; 243 C; 206 G; 223 T; 0 other;

Query Match 79.0%; Score 15.8; DB 20; Length 900;
 Best Local Similarity 89.5%; Pred. No. 34;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 tcacgtgacacagtcacaac 20
 |||| |||||
 Db 176 tcaccgcacacagtcacaac 194

RESULT 13

AAAF0993
 ID AAA0993 standard; DNA; 900 BP.

AC AA09093;
 XX

DT 12-FEB-2001 (first entry)
 XX

DE Human alpha3(IV)NC1 coding sequence.
 XX

KW Type IV collagen alpha chain monomer; human; inhibitor; angiogenesis;
 KW tumour growth; integrin receptor; carcinoma; sarcoma; rhabdomyosarcoma;
 KW retinoblastoma; Bwing sarcoma; neuroblastoma; osteosarcoma; leukaemia;
 KW diabetic retinopathy; rheumatoid arthritis; neovascularisation;
 KW muscular degeneration; corneal graft rejection; vitamin A deficiency;
 KW atopic keratitis; Mycobacteria infection; chemical burn; sarcoid;
 KW Kaposi's sarcoma; sickle cell anaemia; carotid obstructive disease;
 KW chronic inflammation; psoriasis; therapy; alpha3(IV)NC1; ds.

OS Homo sapiens.
 XX

PH Key Location/Qualifiers
 FT COS 40..846
 FT /*tag= a
 FT /product= alpha3(IV)NC1

PN W0200059532-A1.
 XX
 PD 12-OCT-2000.
 XX

PE 31-MAR-2000; 2000WO-0508678.
 XX

PR 01-APR-1999; 99US-0127391.
 XX

PA (BIOS-) BIOTRANUM INC.
 XX

PI Brooks P, Hudson B;
 XX

XX WPI; 2000-664962/64.
 XX

DR P-PSDB; AAY7555.
 XX

PT Use of antagonists of specific integrin receptors for inhibiting
 PT angiogenesis, tumour growth or metastases, or endothelial cell
 PT interactions with the extracellular matrix
 PT
 PS Disclosures: Fig 17c; 78pp; English.

CC This sequence encodes a human type IV collagen alpha chain monomer,
 CC designated alpha3(IV)NC1. The invention relates to a method for
 CC inhibiting angiogenesis, tumour growth or metastases, or endothelial cell
 CC interactions with the extracellular matrix, comprising contacting the
 CC cells or tissue with a polypeptide composition containing antagonists of
 CC specific integrin receptors. The methods and the antagonists are useful
 CC for inhibiting angiogenesis, tumour growth or metastases, or endothelial
 CC cell interaction with the extracellular matrix. The antagonists are also
 CC useful for treating diseases and conditions with accompanying undesired
 CC angiogenesis, e.g. solid and blood-borne tumours (e.g. melanomas,
 CC carcinomas, sarcomas, rhabdomyosarcoma, retinoblastoma, Bwing sarcoma,
 CC neuroblastoma, osteosarcoma or leukaemia). These are also applicable to
 CC treating non-tumorigenic diseases and conditions with accompanying
 CC undesired angiogenesis, e.g. diabetic retinopathy, rheumatoid arthritis,
 CC retinal neovascularisation, choroidal neovascularisation, muscular
 CC degeneration, corneal graft rejection, vitamin A deficiency, atopic
 CC keratitis, Mycobacteria infections, chemical burns, Kaposi's sarcoma,
 CC sickle cell anaemia, sarcoid, carotid obstructive disease, post-laser
 CC complications, chronic inflammation or psoriasis.

SQ Sequence 900 BP; 228 A; 243 C; 206 G; 223 T; 0 other;

Query Match 79.0%; Score 15.8; DB 21; Length 900;
 Best Local Similarity 89.5%; Pred. No. 34;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 tcacgtgacacagtcacaac 20
 |||| |||||
 Db 176 tcaccgcacacagtcacaac 194

RESULT 14

AAAF13921/C
 ID AAFL13921 standard; cDNA; 1276 BP.

AC AAFL13921;
 XX

DT 13-MAR-2001 (first entry)
 XX

DE Aspergillus oryzae EST SEQ ID NO:6444.
 XX

KW Multiple gene expression; filamentous fungal cell; EST;
 KW expressed sequence tag; Fusarium venenatum; Aspergillus niger;
 KW Aspergillus oryzae; Trichoderma reesei; identification; recombination;
 KW culture condition; environmental stress; spore morphogenesis;
 KW metabolic pathway engineering; catabolic pathway engineering; ss.

XX OS Aspergillus oryzae.
XX XY WO2000056762-A2.
XX PN
XX PD 28-SEP-2000.
XX PF 22-MAR-2000; 2000MO-US07781.
XX PR 22-MAR-1999; 99US-0273623.
XX PA (NOVO) NOVO NORDISK BIOTECH INC.
XX PA (NOVO) NOVO NORDISK AS.
XX PI Berka RM, Rey MM, Shuster JR, Kauppinen S, Clausen IG, Olsen PB;
XX DR WPI: 2000-594572/56.
XX PT Monitoring differential expression of genes in filamentous fungal cells
XX PT uses fluorescence-labeled nucleic acids isolated from the cells and a
XX PT substrate of expressed sequence tags -
XX PS Claim 88: Page 2643-2644; 3161pp; English.
XX CC The present invention describes a method for monitoring differential
XX CC expression of genes in a first filamentous fungal (FF) cell relative to
XX CC expression of the same genes in one or more second filamentous fungal
XX CC cells. The method uses fluorescence-labeled nucleic acids isolated from
XX CC the FF cells and a substrate of expressed sequence tags (EST). The ESTs
XX CC are used in the methods for monitoring differential expression of genes
XX CC in a first filamentous fungal (FF) cell relative to expression of the
XX CC same genes in one or more second filamentous fungal cells. Monitoring
XX CC the global expression of genes from FF cells allows the production
XX CC potential of the microorganisms to be improved. New genes may be
XX CC discovered, possible functions of unknown open reading frames can be
XX CC identified and gene copy number variation and stability can be
XX CC monitored. The expression of genes can be used to study how FF cells
XX CC adapt to changes in culture conditions, environmental stress, spore
XX CC morphogenesis, recombination, metabolic or catabolic pathway
XX CC engineering. Using ESTs provides several advantages over genomic or
XX CC random cDNA clones including elimination of redundancy as one spot on an
XX CC array equals one gene or open reading frame, and organization of the
XX CC microarrays based on function of the gene products to facilitate
XX CC analysis of the results. AA070478 to AA071247 represents ESTs from
XX CC Fusarium venenatum; AA071248 to AA071853 represents ESTs from Aspergillus
XX CC niger; AA071854 to AA071878 represents ESTs from Aspergillus oryzae; and
XX CC AA071879 to AA071937 represents ESTs from Trichoderma reesei, which are
XX CC all specifically claimed in the present invention.
XX SS Sequence 1276 BP; 311 A; 319 C; 314 G; 332 T; 0 other;

Query Match 76.0%; Score 15.2; DB 21; Length 1276;
Best Local Similarity 85.0%; Pred. NO. 73;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

CY 1 gtcaagtcacacgtcaaac 20
DB 746 gctctgacacgtcacac 727

RESULT 15
AAC35997
ID AAC35997 standard; DNM; 1801 BP.
XX AAC35997;
XX AC
XX AC AAC35997;
XX DE 17-OCT-2000 (first entry)
XX DE Arabidopsis thaliana DNA fragment SEQ ID NO: 12116.
XX KM Hybridisation assay; genetic mapping; gene expression control;
XX KM protein identification; signal transduction pathway;

KW metabolic pathway; promoter; termination sequence; ss.
XX Arabidopsis thaliana.
XX OS
XX XY EP1033405-A2.
XX PN
XX PD 06-SEP-2000.
XX PF 25-FEB-2000; 2000EP-0301439.
XX PR 25-FEB-1999; 99US-0121825.
XX PR 05-MAR-1999; 99US-0123180.
XX PR 09-MAR-1999; 99US-0123548.
XX PR 23-MAR-1999; 99US-0125788.
XX PR 25-MAR-1999; 99US-0126264.
XX PR 29-MAR-1999; 99US-0126785.
XX PR 01-APR-1999; 99US-0127462.
XX PR 06-APR-1999; 99US-0128234.
XX PR 08-APR-1999; 99US-0128714.
XX PR 16-APR-1999; 99US-0129845.
XX PR 19-APR-1999; 99US-0130077.
XX PR 21-APR-1999; 99US-0130449.
XX PR 23-APR-1999; 99US-0130510.
XX PR 23-APR-1999; 99US-0130891.
XX PR 28-APR-1999; 99US-0131449.
XX PR 30-APR-1999; 99US-0132048.
XX PR 30-APR-1999; 99US-0132407.
XX PR 04-MAY-1999; 99US-0132484.
XX PR 05-MAY-1999; 99US-0132485.
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XX PR 19-MAY-1999; 99US-0134941.
XX PR 20-MAY-1999; 99US-0135124.
XX PR 21-MAY-1999; 99US-0135353.
XX PR 24-MAY-1999; 99US-0135629.
XX PR 25-MAY-1999; 99US-0136021.
XX PR 27-MAY-1999; 99US-0136392.
XX PR 28-MAY-1999; 99US-0136782.
XX PR 01-JUN-1999; 99US-0137222.
XX PR 03-JUN-1999; 99US-0137528.
XX PR 04-JUN-1999; 99US-0137502.
XX PR 07-JUN-1999; 99US-0137724.
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XX PR 18-JUN-1999; 99US-0139500.
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XX PR 18-JUN-1999; 99US-0139502.
XX PR 18-JUN-1999; 99US-0139503.
XX PR 18-JUN-1999; 99US-0139504.
XX PR 18-JUN-1999; 99US-0139505.
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XX PR 18-JUN-1999; 99US-0139512.
XX PR 18-JUN-1999; 99US-0139513.
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XX PR 18-JUN-1999; 99US-0139536.
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XX PR 18-JUN-1999; 99US-0139538.
XX PR 18-JUN-1999; 99US-0139539.
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Job time: 4678 sec

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:18:52 ; Search time 117.39 seconds
(without alignments)
31.563 Million cell updates/sec

Title: US-09-142-095-3
Perfect score: 20
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Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 317530 seqs, 92630169 residues

Total number of hits satisfying chosen parameters: 635060

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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5	14.8	74.0	1990	4	US-09-124-698-154
6	14.8	74.0	1990	4	US-09-127-480-154
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33	13.6	68.0	30	1	US-08-483-554B-40	Sequence 40, Appl
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ALIGNMENTS

RESULT 1
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Sequence 3, Application US/08857464
Patent No. 606450
GENERAL INFORMATION:
APPLICANT: Ralston, Stuart H.
APPLICANT: Grant, Stuart F.A.
TITLE OF INVENTION: Diagnostic and Therapeutic Methods and
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Avenue, N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/857,464
FILING DATE: 16-MAY-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9610281.9
FILING DATE: 16-MAY-1996
ATTORNEY/AGENT INFORMATION:
NAME: Esmond, Robert W.
REGISTRATION NUMBER: 32,893
REFERENCE/DOCKET NUMBER: 1581.0180000
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 22 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: both
MOLECULE TYPE: CDNA
US-08-857-464-3

Query Match 82.0% Score 16.4; DB 3; Length 22;
Best Local Similarity 94.4% Pred. No. 4.2;
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Db 19 TCACGTGACACTCAAA 2

RESULT 2

US-08-991-789A-270/C
Sequence 270, Application US/08991789A
Patent No. 6225054

GENERAL INFORMATION:
APPLICANT: Fridakis, Tony N.
Smith, John M.
Reed, Steven G.

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE
TREATMENT AND DIAGNOSIS OF BREAST CANCER

NUMBER OF SEQUENCES: 292

CORRESPONDENCE ADDRESS:
ADDRESSEE: Seed IP Law Group
STREET: 701 Fifth Avenue, Suite 6300
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104-7092

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/991,789A
FILING DATE: 11-Dec-1997
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Potler, Jane E. R.
REGISTRATION NUMBER: 33,332
REFERENCE/DOCKET NUMBER: 210121.419C3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 270:
SEQUENCE CHARACTERISTICS:
LENGTH: 519 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 270:
US-08-991-789A-270

Query Match 74.0%; Score 14.8; DB 4; Length 519;
Best Local Similarity 88.9%; Pred. No. 41;

Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaa 19

Db 74 TCACGTGACTCAGCCAAA 57

RESULT 3

US-08-967-101-154
Sequence 154, Application US/08967101
Patent No. 5840540

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E

TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts

COUNTRY: U.S.A.

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 154:
SEQUENCE CHARACTERISTICS:

LENGTH: 1990 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: cDNA
US-08-967-101-154

Query Match 74.0%; Score 14.8; DB 2; Length 1990;
Best Local Similarity 84.2%; Pred. No. 49;

Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaa 20

Db 168 TCACGTGACCCAGWAAAC 186

RESULT 4

US-08-592-541-154
Sequence 154, Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E

TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 154:
SEQUENCE CHARACTERISTICS:

LENGTH: 1990 base pairs

```

;      TYPE: nucleic acid
;      STRANDEDNESS: single
;      TOPOLOGY: linear
;      MOLECULE TYPE: CDNA
;
US-08-552-541-154

```

Query Match	74.08;	Score 14.8;	DB 2;	Length 1990;
Best Local Similarity	84.28;	Pred. No. 49;		
Matches 16; Conservative	0;	Mismatches 3;	Indels 0;	Gaps 0;

```

OY      2  tcaagtgacacaggtcaaac 20
          ||| ||||| ||| |||||
Db     168 TCAAGTGACCCAGNCAAC 186

```

RESULT 5
 US-09-124-698-154
 Sequence 154, Application US/09124698
 Patent No. 6117978
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSTOP, PETER H
 APPLICANT: ROMKENS, JOHANNA M
 APPLICANT: FRASER, PAUL E
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 183
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: TESTA, HUMINTZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 Zip: 02110
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC Compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION NUMBER: US/09/124,698

```

Query Match      74.0%; Score 14.8; DB 4; Length 1990;
Best local similarity 84.2%; Pred. No. 49;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0
QY      2 tcaagtcacacagtcacac 20
          ||| ||||| ||||| |||||
Db      168 tcaatgacccacacacaaac 186

```

RESULT 6
US-09-127-480-154
Sequence 154, Application US/09127480

```

: Patent No.6194153
: GENERAL INFORMATION:
: APPLICANT: ST. GEORGE-HYSLOP, PETER H
: APPLICANT: ROMMENS, JOHANNA M
: APPLICANT: FRASER, PAUL E
: TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
: TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
: NUMBER OF SEQUENCES: 183
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: TESTA, HURWITZ & THIBEAULT
: STREET: High Street Tower - 125 High Street
: CITY: Boston
: STATE: Massachusetts
: COUNTRY: U.S.A.
: ZIP: 02110
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/127,480
: FILING DATE:
: CLASSIFICATION:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US/08/592,541
: FILING DATE:
: ATTORNEY/AGENT INFORMATION:
: NAME: Pitcher, Edmund R.
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (617) 248-7000
: TELEFAX: (617) 248-7100
: INFORMATION FOR SEQ. ID NO: 154:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1990 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: US-09-127-480-154
:
: Query Match 74.0%; Score 14.8; DB 4; Length 1990;
: Best Local Similarity 84.2%; Pred. No. 49;
: Matches 16; Conservative 0; Mismatches 3; Indels 0; Caps 0;
:
: 2 taactgacacactcaaac 20
: ||| ||||| ||| |||||
: 168 tcgaatgaccacagcnaac 186
: DB

```

RESULT 7
 US-08-496-841C-154
 Sequence 154, Application US/08496841C
 Patent No. 6210919
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H.
 ROMMENS, JOHANN M.
 FRASER, PAUL E.
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 175
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: DARBY & DARBY, PC
 STREET: 805 THIRD AVENUE
 CITY: NEW YORK
 STATE: NEW YORK
 COUNTRY: U.S.A.
 ZIP: 10022
 COMPUTER READABLE FORM:
 MEDIUM TYPE: FLOPPY DISK
 COMPUTER: IBM PC COMPATIBLE
 OPERATING SYSTEM: PC-DOS/MS-DOS

SOURCE: Patent Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/496,841C

FILING DATE: 28-Jun-1995

CLASSIFICATION: <Unknown>

ATTORNEY/AGENT INFORMATION:

NAME: Paul F. Fehlner, Ph.D.

REGISTRATION NUMBER: 35,115

TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 527-7700

TELEFAX: (212) 753-6237

INFORMATION FOR SEQ ID NO: 154:

SEQUENCE CHARACTERISTICS:

LENGTH: 1990 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

SEQUENCE DESCRIPTION: SEQ ID NO: 154:

US-08-496-841C-154

Query Match 74.0%; Score 14.8; DB 4; Length 1990;
Best Local Similarity 84.2%; Pred. No. 49;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 tcaagtacacagtcacaa 20

Db 168 TCAGTGACCCAGCAGCAAC 186

RESULT 8
US-08-100-247-4

Sequence 4, Application US/08100247

Patent No. 5571787

GENERAL INFORMATION:

APPLICANT: O'BRIEN, JOHN S.

APPLICANT: KISHIMOTO, YASUO

TITLE OF INVENTION: PROSAPOSIN AS A NEUROTROPHIC FACTOR

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: KNOBBE, MARTENS, OLSON AND BEAR

STREET: 620 NEWPORT CENTER DRIVE SIXTEENTH FLOOR

CITY: NEWPORT BEACH

STATE: CA

COUNTRY: USA

ZIP: 92660

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC Compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/100,247

FILING DATE: 19930730

CLASSIFICATION: 514

ATTORNEY/AGENT INFORMATION:

NAME: Israelsen, Ned A.

REGISTRATION NUMBER: 29,655

REFERENCE/DOCKET NUMBER: O'BRIEN.002A

TELECOMMUNICATION INFORMATION:

TELEPHONE: 619-235-8550

TELEFAX: 619-235-0176

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 2740 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

HYPOTHETICAL: NO

ANTI-SENSE: NO

IMMEDIATE SOURCE:

CLONE: PROSAPOSIN cDNA

US-08-100-247-4

Query Match 74.0%; Score 14.8; DB 1; Length 2740;
Best Local Similarity 88.9%; Pred. No. 51;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcaagtacacagtcacaa 19

Db 1194 TCAGTGACTCAGCCAAA 1211

RESULT 9
US-08-483-146A-4

Sequence 4, Application US/08483146A

Patent No. 5636080

GENERAL INFORMATION:

APPLICANT: O'Brien, John S.

APPLICANT: Kishimoto, Yasuo

TITLE OF INVENTION: PHARMACEUTICAL COMPOSITIONS

TITLE OF INVENTION: COMPRISING PROSAPOSIN AND NEUROTROPHIC PEPTIDES DERIVED

NUMBER OF SEQUENCES: 11

CORRESPONDENCE ADDRESS:

ADDRESSEE: Knobb, Martens, Olson and Bear

STREET: 620 Newport Center Blvd. 16th Floor

CITY: Newport Beach

STATE: CA

COUNTRY: USA

ZIP: 92660

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/483,146A

FILING DATE: 07-JUN-1995

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Israelsen, Ned A.

REGISTRATION NUMBER: 29,655

REFERENCE/DOCKET NUMBER: MYELOS.002DV1

TELECOMMUNICATION INFORMATION:

TELEPHONE: 619-235-8550

TELEFAX: 619-235-0176

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 2740 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-08-483-146A-4

Query Match 74.0%; Score 14.8; DB 1; Length 2740;
Best Local Similarity 88.9%; Pred. No. 51;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcaagtacacagtcacaa 19

Db 1194 TCAGTGACTCAGCCAAA 1211

RESULT 10
US-08-232-513A-5

Sequence 5, Application US/08232513A

Patent No. 5700909
GENERAL INFORMATION:
APPLICANT: O'Brien, John S.
TITLE OF INVENTION: Prosaposin and cytokine-derived peptides
TITLE OF INVENTION: as Therapeutic Agents
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell & Flores LLP
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/232.513A
FILING DATE: 21-APR-1994
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/100,247
FILING DATE: 30-JUL-1993
ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-UD 1643
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 2740 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..2740
OTHER INFORMATION: /label=HmL_Prosaposin
US-08-232-513A-5

Query Match 74.0%; Score 14.8; DB 1; Length 2740;
Best Local Similarity 88.9%; Pred. No. 51;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaa 19
|||||
DB 1194 TCACGTGACTCAGCCAAA 1211

RESULT 11
US-08-484-594A-4
Sequence 4, Application US/08484594A
Patent No. 5714459
GENERAL INFORMATION:
APPLICANT: O'Brien, John S.
APPLICANT: Kishimoto, Yasuo
TITLE OF INVENTION: USE OF PROSAPOSIN AND NEUROTROPHIC PEPTIDES
TITLE OF INVENTION: DERIVED THEREFROM
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Knodbe, Martens, Olson and Bear
STREET: 620 Newport Center Drive, Sixteenth Floor
CITY: Newport Beach
STATE: CA
COUNTRY: USA
ZIP: 92660
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/484,594A
FILING DATE: 07-JUN-1995
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/100,247
FILING DATE: 30-JUL-1993
ATTORNEY/AGENT INFORMATION:
NAME: Israelson, Ned A.
REGISTRATION NUMBER: 29,655
REFERENCE/DOCKET NUMBER: MTELOS.0021V2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619-235-8550
TELEFAX: 619-235-0176
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 2740 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-484-594A-4

Query Match 74.0%; Score 14.8; DB 1; Length 2740;
Best Local Similarity 88.9%; Pred. No. 51;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 tcacgtgacacagtcacaa 19
|||||
DB 1194 TCACGTGACTCAGCCAAA 1211

RESULT 12
US-09-060-756-666/c
Sequence 666, Application US/09060756
Patent No. 6183857
GENERAL INFORMATION:
APPLICANT: Cole, Stewart
APPLICANT: Buchleser-Brosch, Roland
APPLICANT: Gordon, Stephen
APPLICANT: Billault, Alain
TITLE OF INVENTION: METHOD FOR ISOLATING A POLYNUCLEOTIDE OF INTEREST FROM
TITLE OF INVENTION: THE GENOME OF A MYCOBACTERIUM USING A BAC-BASED DNA
TITLE OF INVENTION: LIBRARY APPLICATION TO THE DETECTION OF MYCOBACTERIA
FILE REFERENCE: 3495-0169
CURRENT APPLICATION NUMBER: US/09/060,756
CURRENT FILING DATE: 1998-04-16
NUMBER OF SEQ ID NOS: 743
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 666
LENGTH: 448
TYPE: DNA
ORGANISM: Mycobacterium tuberculosis
FEATURE:
NAME/KEY: unsure
LOCATION: (various positions within the sequence)
OTHER INFORMATION: applicants are uncertain of bases designated as "n"
US-09-060-756-666

Query Match 72.0%; Score 14.4; DB 4; Length 448;
Best Local Similarity 88.2%; Pred. No. 65;
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 acgtgacacagtcacaa 20
|||||
DB 298 ACGGGACAGCTCAAC 282

RESULT 13

US-08-053-131-145/c

Sequence 145, Application US/08053131

Patent No. 5661016

GENERAL INFORMATION:

APPLICANT: Lomborg, Nils

APPLICANT: Kay, Robert M.

TITLE OF INVENTION: Transgenic No. 5661016-Human Animals for

PRODUCING Heterologous Antibodies

NUMBER OF SEQUENCES: 197

CORRESPONDENCE ADDRESS:

ADDRESSEE: Townsend and Townsend Kourile and Crew

STREET: One Market Plaza, Stewart Tower, Suite 200

CITY: San Francisco

STATE: California

COUNTRY: USA

ZIP: 94105

COMPUTER READABLE FORM:

MEDIUM TYPE: floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/053,131

FILING DATE: 26-APR-1993

CLASSIFICATION: 800

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/990,860

FILING DATE: 16-DEC-1992

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/810,279

FILING DATE: 17-DEC-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/853,408

FILING DATE: 18-MAR-1992

ATTORNEY/AGENT INFORMATION:

NAME: Smith, William M.

REGISTRATION NUMBER: 30,223

REFERENCE/DOCKET NUMBER: 14643-9-3

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-326-2400

TELEFAX: 415-326-2422

INFORMATION FOR SEQ ID NO: 145:

SEQUENCE CHARACTERISTICS:

LENGTH: 84 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-053-131-145

Query Match 71.0%; Score 14.2; DB 1; Length 84;

Best Local Similarity 84.2%; Pred. No. 66;

Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 gtccagtgacacagtcacaa 19

DB 36 gtccagtgacacagtcacaa 18

RESULT 14

US-08-645-641-145/c

Sequence 145, Application US/08645641

Patent No. 5719032

GENERAL INFORMATION:

APPLICANT: Lomborg, Nils

APPLICANT: Kay, Robert M.

TITLE OF INVENTION: Transgenic No. 5719032-Human Animals for

PRODUCING Heterologous Antibodies

NUMBER OF SEQUENCES: 150

CORRESPONDENCE ADDRESS:

ADDRESSEE: William M. Smith

STREET: Two Embarcadero Center, 8th Floor

CITY: San Francisco

STATE: California

COUNTRY: USA

ZIP: 94111-3834

COMPUTER READABLE FORM:

MEDIUM TYPE: floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/645,641

FILING DATE: 20-MAY-1996

CLASSIFICATION: 800

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/904,068

FILING DATE: 23-JUN-1992

ATTORNEY/AGENT INFORMATION:

NAME: Smith, William M.

REGISTRATION NUMBER: 30,223

REFERENCE/DOCKET NUMBER: 14643-000913

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-326-2400

TELEFAX: 415-326-2422

INFORMATION FOR SEQ ID NO: 145:

SEQUENCE CHARACTERISTICS:

LENGTH: 84 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-645-641-145

Query Match 71.0%; Score 14.2; DB 1; Length 84;

Best Local Similarity 84.2%; Pred. No. 66;

Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 gtccagtgacacagtcacaa 19

DB 36 gtccagtgacacagtcacaa 18

RESULT 15

US-07-853-408B-145/c

Sequence 145, Application US/07853408B

Patent No. 5789650

GENERAL INFORMATION:

APPLICANT: Lomborg, Nils

APPLICANT: Kay, Robert M.

TITLE OF INVENTION: Transgenic No. 5789650-Human Animals for

PRODUCING Heterologous Antibodies

NUMBER OF SEQUENCES: 150

CORRESPONDENCE ADDRESS:

ADDRESSEE: William M. Smith

STREET: One Market Plaza, Stewart Tower, Suite 2000

CITY: San Francisco

STATE: California

COUNTRY: USA

ZIP: 94105

COMPUTER READABLE FORM:

MEDIUM TYPE: floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/853,408B

FILING DATE: 19920318

CLASSIFICATION: 800

ATTORNEY/AGENT INFORMATION:

NAME: Smith, William M.

REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 14643-9
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-326-2400
TELEFAX: 415-326-2422
INFORMATION FOR SEQ. ID NO.: 145:
SEQUENCE CHARACTERISTICS:
LENGTH: 84 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-07-853-408B-145

Query Match 71.0%; Score 14.2; DB 1; Length 84;
Best Local Similarity 84.2%; Pred. No. 66;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 gtacgctgacacagtcaca 19
|||||
Db 36 gtacgcttaccagtcaca 18

Search completed: July 25, 2001, 05:18:53
Job time: 9216 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 04:54:51 ; Search time 2762.24 Seconds
(without alignments)
68.443 Million cell updates/sec

Title: US-09-142-095-3

Perfect score: 20

Sequence: 1 gtacagtgacacagcgaac 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1022815 segs, 4726426750 residues

Total number of hits satisfying chosen parameters: 20456230

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

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2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
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45: em_esthum11:*
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72: em_esthum38:*
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 256: gb_est176:*
 257: gb_est177:*
 258: gb_est178:*

Pred. No. is the number of results predicted by chance to have a
 score greater than or equal to the score of the result being printed,
 and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	length	DB	ID	Description
C 1	18.4	92.0	737	166	BS306792	BS306792 601101955
C 2	17.4	87.0	238	150	BS310653	BS310653 BS310653
C 3	17.4	87.0	295	16	AI104756	AI104756 EST2144045
C 4	17.4	87.0	438	237	A2015172	A2015172 RPCI-23-
C 5	17.4	87.0	482	233	AQ798101	AQ798101 HS 3156
C 6	17.4	87.0	554	14	AI180191	AI180191 EST223932
C 7	16.8	84.0	458	222	FR0039113	AI126681 Fugu rdb
C 8	16.8	84.0	400	222	FR0039183	AI126681 Fugu rdb
C 9	16.8	84.0	513	222	FR0039084	AI126682 Fugu rdb
C 10	16.8	84.0	539	166	BS3681176	BS3681176 601222008
C 11	16.8	84.0	548	149	BF5000440	BF5000440 AI15369
C 12	16.8	84.0	562	190	W73212	W73212 zds310-r1
C 13	16.8	84.0	804	146	BF232658	BF232658 602023222
C 14	16.8	84.0	835	220	CNS0260P	AI195730 Tetraodon
C 15	16.8	84.0	835	221	CNS0260P	AI240268 Tetraodon
C 16	16.8	84.0	968	220	CNS0260P	AI240324 Tetraodon
C 17	16.8	84.0	1046	172	BF982699	BF982699 602305088
C 18	16.8	84.0	1114	154	BG493558	BG493558 602342444
C 19	16.4	82.0	597	231	AO657757	AO657757 Sheared 1
C 20	16.4	82.0	639	222	FR0007351	I 291161 F. rubripes
C 21	16.4	82.0	837	221	CNS0360P	AI253994 Tetraodon
C 22	16.4	82.0	873	150	BF578047	BF578047 602091897
C 23	16.4	82.0	892	220	CNS02010P	AI175751 Tetraodon
C 24	16.4	82.0	988	219	CNS0110P	AI156479 Anopheles
C 25	16	80.0	971	221	CNS0414P	AI296066 Tetraodon
C 26	16	80.0	981	221	CNS0444P	AI274375 Tetraodon
C 27	15.8	79.0	181	250	AZ841578	AB841578 2M013950P
C 28	15.8	79.0	199	127	BS160957	BS160957 BS160957
C 29	15.8	79.0	253	152	BG318736	BG318736 NXPV_016
C 30	15.8	79.0	264	229	AV329412	AV329412 AV329412
C 31	15.8	79.0	279	109	AV062353	AV062353 AV062353
C 32	15.8	79.0	284	25	AV156238	AV156238 AV156238
C 33	15.8	79.0	287	16	AI111900	AI11900 UI-R-Al-e
C 34	15.8	79.0	303	14	AA957109	AA957109 UI-R-Al-e
C 35	15.8	79.0	311	15	AI040776	AI040776 OX15408-S
C 36	15.8	79.0	327	17	AI228082	AI228082 EST224777
C 37	15.8	79.0	345	22	AI2596772	AI2596772 EST251475
C 38	15.8	79.0	370	224	AO130228	AO130228 HS 3007
C 39	15.8	79.0	375	112	AM142511	AM142511 EST92767
C 40	15.8	79.0	381	114	AM285367	AM285367 UI-R-B12
C 41	15.8	79.0	387	122	AM919078	AM919078 EST25038
C 42	15.8	79.0	404	188	R92042	R92042 yp96a04.i1
C 43	15.8	79.0	404	187	RI17651	RI17651 YG15G08.i1
C 44	15.8	79.0	409	14	AA944486	AA944486 EST919965
C 45	15.8	79.0	420	12	AA800910	AA800910 EST910407

ALIGNMENTS

Email: cgapbs-remail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing Arrayed by: Incyte Genomics, Inc.
Clone distribution: MCC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLM8543 row: c column: 13
High quality sequence stop: 359.

FEATURES

source

1..737

/organism="Mus musculus"
/strain="C57BL/6J (f1eral)"
/db_xref="taxon:10090"
/clone="IMAGE:3494340"
/clone_1b="NCI-CGAP-Lu29"
/tissue_type="spontaneous tumor, metastatic to mammary."
stem_cell_origin=""
/lab_host="DH10B"
/note="Organ: lung; Vector: pCMV-SPORT6; Site:1; Salt; Site_2: Not; Cloned unidirectionally. Primer: Oligo dr. Library constructed by Life Technologies. Investigator providing samples: Gilbert Smith, NIH"

BASE COUNT

169 a 190 c 177 g 201 t

ORIGIN

Query Match 92.0%; Score 18.4; DB 166; Length 737;
Best Local Similarity 95.0%; Pred. No. 31;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Caps 0;

Cy 1 gtcaagtgacacagctcaaac 20
|||||
Db 305 gtccagctgacacagctcacc 286

RESULT 2

BB310653/c

LOCUS BB310653 238 bp mRNA EST 10-JUL-2000

DEFINITION BB310653 RIKEN full-length enriched, adult male corpora quadrigemina Mus musculus CDNA clone B230316R05 3', mRNA sequence.

ACCESSION BB310653

VERSION BB310653.1 GI:9011358

KEYWORDS EST

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 238)

REFERENCE 1. Fodor T., Erdos S., Kukukishi Y., Hara A., Havasov N.

URI: <http://genome.rtc.riken.go.jp/>
 Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagao, S., Sasaki, N., Okazaki, Y., Muramatsu, K. and Hayashizaki, Y.
 Thermostabilization and thermocatalytic activation of thermostable enzymes by trehalose and its application for the synthesis of full length cDNA. *Proc. Natl. Acad. Sci. U.S.A.* 95 (2), 520-524 (1998)
 Itoh, M., Katsunari, T., Akiyama, J., Shibata, K., Iwano, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, K., Okazaki, Y. and Hayashizaki, Y.
 Automated filtration-based high-throughput plasmid preparation system. *Genome Res.* 9 (5), 463-470 (1999)
 Carninci, P. and Hayashizaki, Y.
 High-efficiency full-length cDNA cloning. *Methods Enzymol.* 303, 19-44 (1999)
 Please visit our web site (<http://genome.rtc.riken.go.jp/>) for further details.

FEATURES

source

Location/Qualifiers
 1. 238
 /organism="Mus musculus"
 /db_xref="taxon:10090"
 /clone="B230316K05"
 /clone_1id="RIKEN full-length enriched, adult male corpora quadrigemina"
 /sex="male"
 /tissue_type="corpora quadrigemina"
 /dev_stage="adult"
 /lab_host="DH10B"

/note="Site 1: Salt; Site 2: BamHI. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN, Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5' GAGGAGAGAGATCCAGAGCTCTTTTCTTTTCTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot - 20.0 and subtraction to Rot - 459.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGGAGAGATCTTCAGATTAATTAATCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified plasmid pUC19 (+) after bulk excision from Lambda FLX I."

BASE COUNT

71 a 43 c 51 g 73 t

ORIGIN

Query Match

Best Local Similarity 87.0%; Score 17.4; DB 130; Length 238;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 gtcacgtgacacgctcaaa 19
 |||||||
 Db 72 gtcacgtgacacgctcaaa 54

RESULT 3

A1104756

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

295 bp mRNA EST 20-JAN-1999
 EST211045 Normalized rat heart, Bento Soares Rattus sp. cDNA clone
 A1104756
 A1104756.1 GI:3709041
 EST.
 Rattus sp.
 Rattus sp.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.
 1 (bases 1 to 295)
 Lee, N.H., Glodex, A., Chandra, I., Mason, T.M., Quackenbush, J.,
 Keriavage, A.R. and Adams, M.D.

TITLE

Rat Genome Project: Generation of a Rat EST (RST) Catalog & Rat
 Gene Index
 Unpublished (1998)

JOURNAL

COMMENT

Contact: Lee, NH
 The Institute for Genomic Research
 9712, Medical Center Drive, Rockville, MD 20850, USA
 Tel: (301)-838-3529
 Fax: (301)-838-0208
 Email: nhlee@tigr.org
 Seq primer: M13-21.

FEATURES

source

Location/Qualifiers
 1. 295
 /organism="Rattus sp."
 /db_xref="ATCC (Inhost):2025643"
 /db_xref="taxon:10118"
 /clone="RHEC160"
 /clone_1id="Normalized rat heart, Bento Soares"
 /note="Organ: heart; Vector: pT73Pac; Site 1: EcoRI;
 Site 2: NotI"
 BASE COUNT 87 a 49 c 74 g 85 t
 ORIGIN

Query Match 87.0%; Score 17.4; DB 16; Length 295;
 Best Local Similarity 94.7%; Pred. No. 92;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 tcaacgtgacacgctcaac 20
 |||||||
 Db 233 tcaacgtgacacgctcaac 251

RESULT 4

A2015172

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

A2015172 438 bp DNA GSS 25-FEB-2000
 RPCI-23-289M15.TJ RPCI-23 Mus musculus genomic clone RPCI-23-289M15
 , DNA sequence.
 A2015172
 A2015172.1 GI:7090556
 GSS.
 house mouse.
 Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 438)
 Zhao, S., Nierman, M., Feldblum, T., Malek, J., Shatman, S., Aklure, B., Levins, M., McGann, S., Tsegaye, G., Geer, K., Krol, M., de Jong, P. and Fraser, C.M.
 Mouse BAC End Sequences from Library RPCI-23
 Unpublished (1999)
 Other GSSs: RPCI-23-289M15.TV
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the mouse BAC library RPCI-23. For BAC
 library availability, please contact Pieter de Jong
 (pleter@tigr.org, med.bufileo.edu). Clones may be purchased from
 BACPAC Resources (<http://bacpac.med.bufileo.edu/orderingframe.htm>)
 or from Resea ch Genetics (info@resgen.com). BAC end page:
http://www.tigr.org/tidb/bac_ends/mouse/bac_end_intro.html
 Plate: 289 row: M Column: 15
 Seq primer: SP6
 Class: BAC ends.
 Location/Qualifiers
 1. 438
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="RPCI-23-289M15"

Best Local Similarity: 94.7%; Pred. No. 98;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0

TITLE Direct Submission
Submitted (11-OCT-1999) KRC Human Genome Mapping Project Resource
Centre Hinxton, Cambridges, CB10 1SB, UK
E-mail:

COMMENT

biohelphgmp.mrc.ac.uk
Vector: pbluescript II KS
V_type: phagemid
PRIMER: KS
DESCR:

One pass dye-terminator sequencing of cosmid cloned genomic sequence.

FEATURES

SOURCE

BASE COUNT

67 a 66 c 58 g 64 t 3 others

ORIGIN

Query Match 84.0%; Score 16.8; DB 222; Length 258;
Best Local Similarity 90.0%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 gtcaagtgcacagtcacaac 20
||||| ||||||| |||||
Db 200 GTCACATGACACAGTCAACC 181

RESULT 8

FR0039183/c

LOCUS FR0039183 400 bp DNA 22-OCT-1999
DEFINITION Fugu rubripes GSS sequence, clone 090H03cA5, genomic survey sequence.

ACCESSION

AL126681.1 GI:6108296
KEYWORDS GSS; genome survey sequence.

SOURCE

Takifugu rubripes.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes; Tetraodontidae; Takifugu.

REFERENCE

1 (bases 1 to 400)
Elgar, G., Clark, M.S., Smith, S., Meek, S., Warner, S., Edwards, Y.J.K., Umranta, Y., Williams, G. and Brenner, S.

TITLE

Direct Submission

Submitted (11-OCT-1999) MRC Human Genome Mapping Project Resource Centre, Hinxton, Cambridge, CB10 1SB. UK Email:

COMMENT

biohelphgmp.mrc.ac.uk
Vector: pbluescript II KS
V_type: phagemid
PRIMER: KS

DESCR:

One pass dye-terminator sequencing of cosmid cloned genomic sequence.

FEATURES

SOURCE

BASE COUNT

102 a 102 c 88 g 103 t 5 others

ORIGIN

Query Match 84.0%; Score 16.8; DB 222; Length 400;
Best Local Similarity 90.0%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 gtcaagtgcacagtcacaac 20
||||| ||||||| |||||
Db 296 GTCACATGACACAGTCAACC 277

RESULT 9

FR0039084/c
LOCUS FR0039084 513 bp DNA 22-OCT-1999
DEFINITION Fugu rubripes GSS sequence, clone 090H03cA5, genomic survey sequence.

ACCESSION

AL126582.1 GI:6108197

KEYWORDS

GSS; genome survey sequence.

SOURCE

Takifugu rubripes.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes; Tetraodontidae; Takifugu.

REFERENCE

1 (bases 1 to 513)

AUTHORS

Elgar, G., Clark, M.S., Smith, S., Meek, S., Warner, S., Edwards, Y.J.K., Umranta, Y., Williams, G. and Brenner, S.

TITLE

Direct Submission

Submitted (11-OCT-1999) MRC Human Genome Mapping Project Resource Centre, Hinxton, Cambridge, CB10 1SB. UK Email:

COMMENT

biohelphgmp.mrc.ac.uk
Vector: pbluescript II KS
V_type: phagemid
PRIMER: KS

DESCR:

One pass dye-terminator sequencing of cosmid cloned genomic sequence.

FEATURES

SOURCE

BASE COUNT

132 a 130 c 112 g 134 t 5 others

ORIGIN

Query Match 84.0%; Score 16.8; DB 222; Length 513;
Best Local Similarity 90.0%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 gtcaagtgcacagtcacaac 20
||||| ||||||| |||||
Db 30 GTCACATGACACAGTCAACC 11

RESULT 10

BE368176

LOCUS BE368176 539 bp mRNA 21-JUL-2000
DEFINITION 601222089F1 NCI_CGAP_Lu29 Mus musculus cDNA clone IMAGE:359711 5', mRNA sequence.

ACCESSION

BE368176 GI:9313539

VERSION

EST.

KEYWORDS

EST.

SOURCE

house mouse.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

1 (bases 1 to 539)

AUTHORS

NH MGC. <http://mgc.ncl.nih.gov/>.

TITLE

National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL

Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs@ncl.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: L14M8759 row: b column: 24
High quality sequence stop: 536.

FEATURES

Location/Qualifiers

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/organism="Mus musculus"
/strain="C57BL/6J (fetal)"
/db_xref="taxon:10090"
/clone_image="3590711"
/clone_id="NCI_CGAP_Lu29"
/tissue_type="spontaneous tumor, metastatic to mammary.
stem cell origin."
/lab_host="DH10B"
/notes="Organ: lung; Vector: pCMV-SPORT6; Site: 1; Site: 2; Note: Cloned unidirectionally. Primer: Oligo dr.
Library constructed by Life Technologies. Investigator
providing samples: Gilbert Smith, NIH"

BASE COUNT
201 a 92 c 143 g 103 t

ORIGIN
Query Match 84.0%; Score 16.8; DB 166; Length 539;
Best Local Similarity 90.0%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

cy 1 gtcactgacacgtcacac 20
|||||
Db 207 gtcactgacacgtcacac 226

RESULT 11
BF500440/c 548 bp mRNA EST 16-APR-2001
LOCUS BF500440.1 GI:11583741
DEFINITION BF500440.1 BF500440
VERSION BF500440.1 GI:11583741
KEYWORDS EST.
SOURCE Drosophila melanogaster
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 548)
REFERENCE
1 Stapleton, M., Brokstein, P., Hong, L., Agdayani, A., Baxter, E., Berman,
B., Carlson, J., Champe, M., Chavez, C., Chew, M., Dorsett, V., Farfan,
D., Frise, E., George, R., Gonzalez, M., Guarin, H., Harris, N., Li, P.,
Liao, G., Miranda, A., Misra, S., Mungall, C. J., Nuno, J., Pacleb, J.,
Park, S., Paragás, A., Phoumenavong, S., Wan, K., Yu, C., Lewis, S. E.,
Celniker, S. and Rubin, G. M.
Berkeley Drosophila Gene Collection Project
Unpublished (2000)
COMMENT
Contact: Stapleton, M.
BDGP
Lawrence Berkeley National Lab
One Cyclotron Rd, Berkeley, CA 94720, USA
Fax: 510 486 6798
Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
hit genomic sequence AEO03606
Plate: AT.153 row: F column: 9
High quality sequence strop: 533.
Location/Qualifiers
1. .548
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_image="AT15369"
/clone_id="AT15369"
/clone_host="AT Drosophila melanogaster adult testes POTB7"
/sex="male"
/dest_stage="0-3 day old Ore-R males"
/lab_host="Plates AT.10-AT.120: DHS-alpha. Plates
AT.121-AT.319: DHS-alpha 10nA
/notes="Organ: ADULT testes; Vector: POTB7; Site: 1; Site: 2; Note: The mRNA for the testis library was made
from testes and seminal vesicles and dissected from 0-3
day old Ore-R males. RNA kindly provided by the lab of
Margaret Fuller. Sized fractionated cDNAs were directly

```

[illegible]

QY 1 gtcacgtgacacagtcacac 20
||||| ||||| |||||
DB 549 GTCACGTGACACAGTCACAC 530

RESULT 13
BF232638/c 804 bp mRNA EST 14-NOV-2000
LOCUS 602023221F1 NCI_CGAP_L19 Mus musculus cDNA clone IMAGE:4158796 5'
DEFINITION mRNA sequence.
ACCESSION BF232638
VERSION BF232638.1 GI:11142204
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 804)
AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE NIH-MGC http://mgc.ncl.nih.gov/
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT Unpublished (1999).
Contact: Robert Strausberg, Ph.D.
Email: cgsb@ncl.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LMA9435 row: 1 column: 05
High quality sequence stop: 657.

FEATURES
source
1. 804
Location/Qualifiers
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone IMAGE:4158796"
/clone_11b="NCI_CGAP_L19"
/lab_host="DH10B (T1 phage-resistant)"
/note="Organ: liver; Vector: pCMV-SPORT6; Site: 1: NCI; Site: 2: Salt; cloned unidirectionally. Primer: Oligo dT. Average insert size 1.9 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP Library."

BASE COUNT 196 a 216 c 233 g 159 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 146; Length 804;
Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 gtcacgtgacacagtcacac 20
||||| ||||| |||||
DB 455 GTCACGTGACACAGTCACAC 436

RESULT 14
CNS0260P/c 835 bp DNA GSS 13-MAY-2000
LOCUS Tetradon nigriviridis genome survey sequence T7 end of clone
DEFINITION 139P15 of library G from Tetradon nigriviridis, genomic survey sequence.
ACCESSION AL195730
VERSION AL195730.1 GI:7833880
KEYWORDS GSS; genome survey sequence.
SOURCE Tetradon nigriviridis.
ORGANISM Tetradon nigriviridis
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percormorpha; Tetraodontiformes; Tetraodontidae; Tetradon.

REFERENCE 1 (bases 1 to 835)
AUTHORS Roest-Crollius, H., Jallion, O., Dasilva, C., Fizesmes, C., Fisher, C., Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and Weissenbach, J.
TITLE Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigriviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 835)
AUTHORS Roest-Crollius, H., Jallion, O., Dasilva, C., Bouneau, L., Fisher, C., Bernot, A., Fizesmes, C., Winkler, P., Brotter, P., Quetier, F., Saurin, W. and Weissenbach, J.
TITLE Human gene number estimate provided by genome wide analysis using Tetradon nigriviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 835)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/Genbank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetradon nigriviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/tetradon>.

FEATURES
source
1. 835
Location/Qualifiers
/organism="Tetradon nigriviridis"
/db_xref="taxon:99883"
/clone="135P15"
/clone_11b="G"
/note="Genoscope sequence ID : COA0135CH08UP1-end : T7"

BASE COUNT 192 a 195 c 246 g 200 t 2 others
ORIGIN

Query Match 84.0%; Score 16.8; DB 220; Length 835;
Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 gtcacgtgacacagtcacac 20
||||| ||||| |||||
DB 531 GTCACGTGACACAGTCACAC 512

RESULT 15
CNS03EDV 835 bp DNA GSS 17-MAY-2000
LOCUS Tetradon nigriviridis genome survey sequence pUC-ori end of clone
DEFINITION 020619 of library G from Tetradon nigriviridis, genomic survey sequence.
ACCESSION AL240288
VERSION AL240288.1 GI:7961037
KEYWORDS GSS; genome survey sequence.
SOURCE Tetradon nigriviridis.
ORGANISM Tetradon nigriviridis
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percormorpha; Tetraodontiformes; Tetraodontidae; Tetradon.

REFERENCE 1 (bases 1 to 835)
AUTHORS Roest-Crollius, H., Jallion, O., Dasilva, C., Fizesmes, C., Fisher, C., Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and Weissenbach, J.
TITLE Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigriviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 835)
AUTHORS Roest-Crollius, H., Jallion, O., Dasilva, C., Bouneau, L., Fisher, C., Bernot, A., Fizesmes, C., Winkler, P., Brotter, P., Quetier, F., Saurin, W. and Weissenbach, J.
TITLE Human gene number estimate provided by genome wide analysis using Tetradon nigriviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 835)
AUTHORS Genoscope.

TITLE
JOURNAL
COMMENT

Direct Submission
Submitted (12-Apr-2000) to the EMBL/GenBank/DBJ databases
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetradon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetradon>.

FEATURES

source

1..835

location/Qualifiers

/organism="Tetradon nigroviridis"

/db_xref="taxon:99883"

/clone="020G19"

/note="Genoscope sequence ID : C08G020AD10SP1-end ;

PUC-Orig"

BASE COUNT 185 a 245 c 233 g 168 t 4 others

ORIGIN

Query Match

84.0%; Score 16.8; DB 221; Length 835;

Best Local Similarity 90.0%; Pred. No. 2.2e+02;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 gtcaagtgacacagtcacac 20

Db 261 gtcaagtgacacagtcacac 280

Search completed: July 25, 2001, 04:54:54
Job time: 10427 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OW nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:17:09 ; Search time 1290.33 Seconds
(without alignments)
227.761 Million cell updates/sec

Title: US-09-142-095-4

Perfect score: 19
Sequence: 1 ttgtctctccagagctt 19

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1344157 seqs, 7733874588 residues

Total number of hits satisfying chosen parameters: 268314

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenDbml:*

- 1: gb_da1:*
- 2: gb_da2:*
- 3: gb_da3:*
- 4: gb_da4:*
- 5: gb_da5:*
- 6: gb_da6:*
- 7: gb_da7:*
- 8: gb_da8:*
- 9: gb_da9:*
- 10: gb_da10:*
- 11: gb_da11:*
- 12: gb_da12:*
- 13: gb_da13:*
- 14: gb_da14:*
- 15: gb_da15:*
- 16: gb_da16:*
- 17: gb_da17:*
- 18: gb_da18:*
- 19: gb_da19:*
- 20: gb_da20:*
- 21: gb_da21:*
- 22: gb_da22:*
- 23: gb_da23:*
- 24: gb_da24:*
- 25: gb_da25:*
- 26: gb_da26:*
- 27: gb_da27:*
- 28: gb_da28:*
- 29: gb_da29:*
- 30: gb_da30:*
- 31: gb_da31:*
- 32: gb_da32:*
- 33: gb_da33:*
- 34: gb_da34:*
- 35: gb_da35:*
- 36: gb_da36:*
- 37: gb_da37:*
- 38: gb_da38:*
- 39: gb_da39:*
- 40: gb_da40:*
- 41: gb_da41:*
- 42: gb_da42:*
- 43: gb_da43:*

44: em_ov:*

45: em_pa:*

46: em_ph:*

47: em_pi:*

48: em_po:*

49: em_pt:*

50: em_sy:*

51: em_un:*

52: em_vl:*

53: gb_sts1:*

54: gb_sts2:*

55: gb_sts3:*

56: gb_sy:*

57: gb_un:*

58: gb_vl:*

59: gb_vl2:*

60: gb_vl3:*

61: gb_vl4:*

62: gb_vl5:*

63: gb_vl6:*

64: gb_vl7:*

65: gb_vl8:*

66: gb_vl9:*

67: gb_vl10:*

68: gb_vl11:*

69: gb_vl12:*

70: gb_vl13:*

71: gb_vl14:*

72: gb_vl15:*

73: gb_vl16:*

74: gb_vl17:*

75: gb_vl18:*

76: gb_vl19:*

77: gb_vl20:*

78: gb_vl21:*

79: gb_vl22:*

80: gb_vl23:*

81: gb_vl24:*

82: gb_vl25:*

83: gb_vl26:*

84: gb_vl27:*

85: gb_vl28:*

86: gb_vl29:*

87: gb_vl30:*

88: gb_vl31:*

89: gb_vl32:*

90: gb_vl33:*

91: gb_vl34:*

92: gb_vl35:*

93: gb_vl36:*

94: gb_vl37:*

95: gb_vl38:*

96: gb_vl39:*

97: gb_vl40:*

98: gb_vl41:*

pred. NO. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result NO.	Score	Query Match Length	ID	Description
1	19	100.0	9	AF135465 Sequence 4
2	19	100.0	89	AF135465 Gorilla g
3	19	100.0	200	AF135466 Pongo pyg
4	19	100.0	207	AF135470 Cebus ape
5	19	100.0	208	AF135464 Gorilla g
6	19	100.0	208	AF135471 Salimiri b
7	19	100.0	234	AF135462 Pan panis
8	19	100.0	89	AF135463 Pan trogl

REFERENCE 2 (bases 1 to 200)
 AUTHORS Ydazeta,G., Hall,D. and Di Rienzo,A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
 SOURCE 1..200
 /organism="Pongo pygmaeus"
 /db_xref="taxon:9600"
 mRNA <121..>200
 /gene="UGT1A1"
 /product="UDP-glucuronosyltransferase 1A1"
 <121..>200
 /gene="UGT1A1"
 121..>200
 /gene="UGT1A1"
 /codon_start=1
 /product="UDP-glucuronosyltransferase 1A1"
 /protein_id="AA09177.1"
 /db_xref="GI:6456550"
 /translation="WAVESQGGPVLVGLLCVLPVYSHA"
 CDS
 BASE COUNT 40 a 46 c 62 g 52 t
 ORIGIN

Query Match 100.0%; Score 19; DB 89; Length 200;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgtctctgcagaggtt 19
 |||||
 Db 114 TTGCTCTGCCAGAGTT 96

RESULT 4
 AF135470/c 207 bp DNA PRI 21-NOV-1999
 LOCUS Cebus apella UDP-glucuronosyltransferase 1A1 (UGT1A1) gene.
 DEFINITION Promoter region and partial cds.
 ACCESSION AF135470
 VERSION AF135470.1 GI:6456557
 KEYWORDS brown capuchin.
 SOURCE Cebus apella
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Cebinae; Cebus.
 1 (bases 1 to 207)
 Hall,D., Ydazeta,G., Destro-Bisol,G., Petzl-Erler,M.L. and Di
 Rienzo,A.
 TITLE Variability at the uridine diphosphate glucuronosyltransferase 1A1
 JOURNAL Pharmacogenetics (1999) In press
 AUTHORS 2 (bases 1 to 207)
 Ydazeta,G., Hall,D. and Di Rienzo,A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
 SOURCE 1..207
 /organism="Cebus apella"
 /db_xref="taxon:9515"
 mRNA <144..>207
 /gene="UGT1A1"
 /product="UDP-glucuronosyltransferase 1A1"
 <144..>207
 /gene="UGT1A1"
 144..>207
 /gene="UGT1A1"
 /codon_start=1
 /product="UDP-glucuronosyltransferase 1A1"
 /protein_id="AA09181.1"
 /db_xref="GI:6456558"
 /translation="VPAQCPAAVCAVCAQPSVPCWE"

BASE COUNT 43 a 46 c 62 g 56 t
 ORIGIN

Query Match 100.0%; Score 19; DB 89; Length 207;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgtctctgcagaggtt 19
 |||||
 Db 114 TTGCTCTGCCAGAGTT 96

RESULT 5
 AF135464/c 208 bp DNA PRI 21-NOV-1999
 LOCUS Gorilla gorilla UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
 DEFINITION promoter region and partial cds.
 ACCESSION AF135464
 VERSION AF135464.1 GI:6456545
 KEYWORDS gorilla.
 SOURCE Gorilla gorilla
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Gorilla.
 1 (bases 1 to 208)
 Hall,D., Ydazeta,G., Destro-Bisol,G., Petzl-Erler,M.L. and Di
 Rienzo,A.
 TITLE Variability at the uridine diphosphate glucuronosyltransferase 1A1
 JOURNAL Pharmacogenetics (1999) In press
 AUTHORS 2 (bases 1 to 208)
 Ydazeta,G., Hall,D. and Di Rienzo,A.
 TITLE Direct Submission
 JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
 SOURCE 1..208
 /organism="Gorilla gorilla"
 /db_xref="taxon:9593"
 /note="from Yerkes Regional Primate Center"
 mRNA <133..>208
 /gene="UGT1A1"
 /product="UDP-glucuronosyltransferase 1A1"
 <133..>208
 /gene="UGT1A1"
 133..>208
 /gene="UGT1A1"
 /codon_start=1
 /product="UDP-glucuronosyltransferase 1A1"
 /protein_id="AA09175.1"
 /db_xref="GI:6456546"
 /translation="WAVESQGGPVLVGLLCVLPVYSHA"

BASE COUNT 43 a 47 c 62 g 56 t
 ORIGIN

Query Match 100.0%; Score 19; DB 89; Length 208;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgtctctgcagaggtt 19
 |||||
 Db 126 TTGCTCTGCCAGAGTT 108

RESULT 6
 AF135471/c 208 bp DNA PRI 21-NOV-1999
 LOCUS Salimti boliviensis UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
 DEFINITION promoter region and partial cds.
 ACCESSION AF135471
 VERSION AF135471.1 GI:6456559

KEYWORDS
Bolivian squirrel monkey.

SOURCE
Saimiri boliviensis

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Cebinae; Saimiri.

REFERENCE
1 (bases 1 to 208)
Hall, D., Ybazaeta, G., Destro-Bisol, G., Petzl-Erler, M.L. and Di Rienzo, A.

TITLE
Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates

JOURNAL
Pharmacogenetics (1999) In press

REFERENCE
2 (bases 1 to 208)
Ybazaeta, G., Hall, D. and Di Rienzo, A.

TITLE
Direct Submission

JOURNAL
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
source
1..208
/organism="Saimiri boliviensis"
/db_xref="taxon:27679"
mRNA
<142..>208
/gene="UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
<142..>208
/gene="UGT1A1"
142..>208
/gene="UGT1A1"
/codon_start=1
/product="UDP-glucuronosyltransferase 1A1"
/protein_id="AA09182.1"
/db_xref="GI:6456560"
/translation="MAASRGCGCVLGLCVLPVSHAK"
BASE COUNT
44 a 52 c 58 g 54 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 19; DB 89; Length 208;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
|||||
Db 135 TTGCTCTGCGCAGAGTT 117

RESULT 7
AF135462/c 234 bp DNA PRI 21-NOV-1999
LOCUS
Pan paniscus UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
promoter region and partial cds.
ACCESSION
AF135462
VERSION
AF135462.1 GI:6456541
KEYWORDS
pygmy chimpanzee.
SOURCE
Pan paniscus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Pan.
REFERENCE
1 (bases 1 to 234)
Hall, D., Ybazaeta, G., Destro-Bisol, G., Petzl-Erler, M.L. and Di Rienzo, A.

TITLE
Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates

JOURNAL
Pharmacogenetics (1999) In press

REFERENCE
2 (bases 1 to 234)
Ybazaeta, G., Hall, D. and Di Rienzo, A.

TITLE
Direct Submission

JOURNAL
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
source
1..234
/organism="Pan paniscus"
/db_xref="taxon:9597"

MRNA
<148..>234
/gene="UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
<148..>234
/gene="UGT1A1"
148..>234
/gene="UGT1A1"
/codon_start=1
/product="UDP-glucuronosyltransferase 1A1"
/protein_id="AA09173.1"
/db_xref="GI:6456542"
/translation="MAVESGCGRPVGLGLCVLPVSHAK"
BASE COUNT
50 a 53 c 69 g 62 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 19; DB 89; Length 234;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
|||||
Db 141 TTGCTCTGCGCAGAGTT 123

RESULT 8
AF135463/c 234 bp DNA PRI 21-NOV-1999
LOCUS
Pan troglodytes UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
promoter region and partial cds.
ACCESSION
AF135463
VERSION
AF135463.1 GI:6456543
KEYWORDS
chimpanzee.
SOURCE
Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Pan.
REFERENCE
1 (bases 1 to 234)
Hall, D., Ybazaeta, G., Destro-Bisol, G., Petzl-Erler, M.L. and Di Rienzo, A.

TITLE
Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates

JOURNAL
Pharmacogenetics (1999) In press

REFERENCE
2 (bases 1 to 234)
Ybazaeta, G., Hall, D. and Di Rienzo, A.

TITLE
Direct Submission

JOURNAL
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
source
1..234
/organism="Pan troglodytes"
/db_xref="taxon:9598"
<148..>234
/gene="UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
<148..>234
/gene="UGT1A1"
148..>234
/gene="UGT1A1"
/codon_start=1
/product="UDP-glucuronosyltransferase 1A1"
/protein_id="AA09174.1"
/db_xref="GI:6456544"
/translation="MAVESGCGRPVGLGLCVLPVSHAK"
BASE COUNT
50 a 53 c 69 g 62 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 19; DB 89; Length 234;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
|||||

Db 141 TTTCCTCTCCGACAGGTT 123

|||||

RESULT 9
AF357220/c 292 bp DNA PRI 27-MAR-2001
LOCUS Homo sapiens bilirubin UDP-glucuronosyltransferase 1 (UGT1) gene,
DEFINITION UGT1*1 allele, promoter and partial cds.
ACCESSION AF357220
VERSION AF357220.1 GI:13448828
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 292)
McKie,K., Addington,T., Nguyen,T.S., Glendenning,M., Kutlar,F. and
Kutlar,A.
TITLE Detection of TATA box TA repeat region [6(TA)repeat] of human
bilirubin UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in an
African American individual
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 292)
McKie,K., Addington,T., Nguyen,T.S., Glendenning,M., Kutlar,F. and
Kutlar,A.
TITLE Direct Submission
REFERENCE Submitted (07-MAR-2001) Medicine/Hemoglobin DNA Laboratory, Medical
AUTHORS College of Georgia, 15th Street, Augusta, GA 30912, USA
JOURNAL Location/Qualifiers
FEATURES
source 1..292
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2"
/map="2q37"
/cell_type="WBC"
/tissue_type="whole blood"
1..>292
/gene="UGT1"
/allele="UGT1*1"
1..64
/gene="UGT1"
28..39
/gene="UGT1"
/note="most common type of repeat; 6(TA)"
28..42
/gene="UGT1"
28..39
/note="polymorphic region"
/rpt_type=tandem
/rpt_unit=ta
65..>292
/gene="UGT1"
/product="bilirubin UDP-glucuronosyltransferase 1"
81..>292
/gene="UGT1"
/codon_start=1
/product="bilirubin UDP-glucuronosyltransferase 1"
/protein_id="AAK27223.1"
/db_xref="GI:13448829"
/translation="MAVESOGGRPLVGLLCLVLPVYSHAKILLIPVDSHMLSL
GAIQOLQORGHETIVLAPDASIVRPG"

BASE COUNT 60 a 69 c 95 g 68 t
ORIGIN

Query Match 100.0%; Score 19; DB 89; Length 292;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctccagaggtt 19
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Db 74 TTTCCTCTCCGACAGGTT 56

RESULT 10
AF352795 531 bp DNA PRI 10-APR-2001
LOCUS Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1A1) gene,
DEFINITION UGT1A1*1 allele, partial cds.
ACCESSION AF352795
VERSION AF352795.1 GI:13569708
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 531)
McKie,K., Kutlar,F., Glendenning,M. and Kutlar,A.
TITLE 7(TA) repeat polymorphism of the TATA box of human bilirubin
UDP-glucuronosyltransferase 1-1(UGT1A1*1) gene in a patient with
sickle cell anemia + high bilirubinemia
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 531)
McKie,K., Kutlar,F., Glendenning,M. and Kutlar,A.
TITLE Direct Submission
REFERENCE Submitted (23-FEB-2001) Medicine/Hemoglobin DNA Laboratory, Medical
AUTHORS College of Georgia, 15th St., AC-1000, Augusta, GA 30912, USA
JOURNAL Location/Qualifiers
FEATURES
source 1..531
/organism="Homo sapiens"
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1..>531
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1..313
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279..295
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279..292
/note="Repeat polymorphism compared to UGT1A1 sequence
presented in GenBank Accession Number AF180372; contains 7
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bilirubinemia"
/rpt_type=tandem
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314..>531
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334..>531
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/codon_start=1
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/protein_id="AAK31204.1"
/db_xref="GI:13569709"
/translation="MAVESOGGRPLVGLLCLVLPVYSHAKILLIPVDSHMLSL
GAIQOLQORGHETIVLAPDASIVRPG"

BASE COUNT 120 a 121 c 137 g 153 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 14;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctccagaggtt 19
|||||
Db 327 TTTCCTCTCCGACAGGTT 309

RESULT 11

AF180372/c 541 bp DNA PRI 05-OCT-1999
 LOCUS Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1) gene,
 UGT1*1 allele, partial cds.
 ACCESSION AF180372
 VERSION AF180372.1 GI:6010649
 KEYWORDS
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 541)
 AUTHORS Kullar,F., Stomak,E., Leitner,C., Nechtman,J. and Kullar,A.
 TITLE Detection of the TATA box polymorphism of the human bilirubin
 UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in a patient with
 sickle cell anemia
 JOURNAL Unpublished

REFERENCE 2 (bases 1 to 541)
 AUTHORS Kullar,F., Stomak,E., Leitner,C., Nechtman,J. and Kullar,A.
 TITLE Direct Submission
 JOURNAL Submitted (24-AUG-1999) Medicine, Hematology/Oncology-Sickle Cell
 Center, Medical College of Georgia, 15th Street, AC-1000, Augusta,
 GA 30912, USA

FEATURES
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 /tissue_type="whole blood"
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/gene="UGT1"
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 1..313

promoter

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 326
 /replace="t"
 279..288

variation

repeat_region

/note="Polymorphic region"
 /rpt_type="tandem"
 /rpt_unit="ca
 279..281

TATA_signal

/gene="UGT1"
 /note="Ritter,J.K., et al., 1992, J. Biol. Chem.,
 267:3257-3261"
 314..>541

mRNA

/gene="UGT1"
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 330..>541

CDS

/gene="UGT1"
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 /codon_start=-1
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 /protein_id="AF01205.1"

BASE COUNT

123 a 124 c 141 g 153 t
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ORIGIN

Query Match 100.0%; Score 19; DB 89; Length 541;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
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 Db 323 TTGCTCCTGCCAGAGGTT 305

RESULT 12

AF180372/c A65504 620 bp DNA PAT 29-MAR-1999
 LOCUS A65504
 DEFINITION Sequence 5 from Patent WO9732042.
 ACCESSION A65504
 VERSION A65504.1 GI:453239

KEYWORDS
 SOURCE unidentified.
 ORGANISM unidentified.
 unclassified.

REFERENCE 1 (bases 1 to 620)
 AUTHORS Burchell,B.
 TITLE DRUG TRIAL ASSAY SYSTEM
 JOURNAL Patent: WO 9732042-A 5 04-SEP-1997;
 UNIT DUNDEE (GB)
 COMMENT Other publication AU 2224197 19970916.
 location/Qualifiers

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BASE COUNT 157 a 127 c 151 g 165 t
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Query Match 100.0%; Score 19; DB 9; Length 620;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
 |||||
 Db 605 TTGCTCCTGCCAGAGGTT 587

RESULT 13

AF110194/c AF110194 918 bp DNA PRI 02-JAN-2001
 LOCUS AF110194
 DEFINITION Homo sapiens chromosome 2 UDP-glucuronosyltransferase (UGT1A1)
 gene, UGT1A1*33 allele, partial cds.

ACCESSION AF110194
 VERSION AF110194.1 GI:12002134
 KEYWORDS
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 918)
 AUTHORS Gullerme,C.
 TITLE Direct Submission
 JOURNAL Submitted (01-DEC-1998) Center for Cancer Research, MIT, 77
 Massachusetts Avenue, E17-540, Cambridge, MA 02139, USA

FEATURES
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 /organism="Homo sapiens"
 /db_xref="taxon:9606"
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 <38..>918
 /note="UGT1A1"
 /note="22..918; 1A1 variant allele; L233R"
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/gene="UGT1A1"
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 /product="UDP-glucuronosyltransferase"
 /protein_id="A654197.1"

CDS

/gene="UGT1A1"
 /codon_start=-1
 /product="UDP-glucuronosyltransferase"
 /protein_id="A654197.1"

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 DSFLQRIKIKYKIKDSAMLISCSHLHNELMASIAESSFVMLTDPFLPCSPIV
 AOVKISLTFVFLHALPRLSPEATQGNPFSPVRLSSHDHMTFLQKKNMLIAFS
 ONFLCQVYSPYATNASEFLQRETVQDLSASVWLFRSDVYVDYRPIIMNVFVG

BASE COUNT 196 a 245 c 235 g 242 t
 ORIGIN

Query Match 100.0%; Score 19; DB 88; Length 918;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcacagagtt 19
 |||
 Db 31 ttgtctctgcacagagtt 13

RESULT 14
 HUMUGT1A 1190 bp DNA PRI 14-JAN-1995
 LOCUS
 DEFINITION Human bilirubin UDP-glucuronosyltransferase (UGT1A) gene isozyme
 exon 1.

ACCESSION M84125 GI:340131
 VERSION
 KEYWORDS bilirubin UDP-glucuronosyltransferase; isozyme.
 SOURCE Homo sapiens (tissue library: cosmid) liver DNA.
 ORGANISM

REFERENCE
 AUTHORS Rittler, J.K., Crawford, J.M. and Owens, I.S.
 TITLE Cloning of two human liver bilirubin UDP-glucuronosyltransferase
 cDNAs with expression in COS-1 cells
 J. Biol. Chem. 266 (2), 1043-1047 (1991)

JOURNAL
 MEDLINE 91093210
 REFERENCE 2 (bases 1 to 1190)
 AUTHORS Rittler, J.K., Chen, F., Sheen, Y.Y., Tran, H.M., Kimura, S.,
 Yeatman, M.T. and Owens, I.S.
 TITLE A novel complex locus UGT1 encodes human bilirubin, phenol, and
 other UDP-glucuronosyltransferase isozymes with identical carboxyl
 termini
 J. Biol. Chem. 267 (5), 3257-3261 (1992)

JOURNAL
 MEDLINE 92147680
 REFERENCE 1 (bases 1 to 1190)
 AUTHORS Rittler, J.K., Chen, F., Sheen, Y.Y., Tran, H.M., Kimura, S.,
 Yeatman, M.T. and Owens, I.S.
 TITLE A novel complex locus UGT1 encodes human bilirubin, phenol, and
 other UDP-glucuronosyltransferase isozymes with identical carboxyl
 termini
 J. Biol. Chem. 267 (5), 3257-3261 (1992)

FEATURES
 source location/Qualifiers
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 /db_xref="taxon:9606"
 /tissue_type="liver"
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 /lab_host="X11-Blue"
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 /number=1
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 85..>948
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 /protein_id="AA61248.1"
 /db_xref="GI:340132"
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 GAIOOLQORHEIVIVADASILYRDAFTLTYPVPEQREDAVESFVMLDPLPCSPY
 DSFLQRIKYKKIKDSAMLSGSHLLHNKELASLESFVMLDPLPCSPY
 AOVLSLPTVFPFLHALPCSLFEATQCPNPSYVPLSLSDPHMTFLQRYKNMLIAPS
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 949..1190
 /partial
 /gene="UGT1A"
 /note="does not fit consensus"
 /number=1

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 TATA_signal
 exon
 gene
 CDS
 intron

BASE COUNT 261 a 290 c 286 g 353 t
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Query Match 100.0%; Score 19; DB 97; Length 1190;
 Best Local Similarity 100.0%; Pred. No. 13;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ttgtctctgcacagagtt 19
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 Db 78 ttgtctctgcacagagtt 60

RESULT 15
 DB67674 3341 bp DNA PRI 14-APR-2000
 LOCUS
 DEFINITION Homo sapiens gene for bilirubin UDP-glucuronosyltransferase 1,
 promoter region and partial cds.
 DB67674
 VERSION DB67674.1 GI:3059176
 KEYWORDS bilirubin UDP-glucuronosyltransferase 1.
 SOURCE Homo sapiens DNA.
 ORGANISM

REFERENCE
 AUTHORS Ueyama, H., Koizumi, O., Soeda, Y., Sato, H., Sato, Y., Okubo, I. and
 Doi, Y.
 TITLE Analysis of the promoter of human bilirubin
 UDP-glucuronosyltransferase gene (UGT1*1) in relevance to Gilbert's
 syndrome
 Hepatol. Res. 9, 152-163 (1997)

JOURNAL
 MEDLINE 91093210
 REFERENCE 2 (bases 1 to 3341)
 AUTHORS Ueyama, H.
 TITLE Direct Submission
 JOURNAL Submitted (04-SEP-1996) to the DDBJ/EMBL/Genbank databases, Hisao
 Ueyama, Shiga University of Medical Science, Department of Medical
 Biochemistry, Seto, Otsu, Shiga 520-21, Japan (Tel:077-548-2162,
 Fax:077-548-2164)
 Sequence updated (08-Jan-1997) by: Hisao Ueyama.

COMMENT
 source location/Qualifiers
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 /db_xref="taxon:9606"
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 2606..2610
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 3088..3097
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 /note="E-box"
 3101..3113
 /note="HNF-1 site"
 3125..3129
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 3149..3153
 TATA_signal
 3177..3341
 /number=1
 3182..3341
 /gene="UGT1*1"
 3192..>3341
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 /codon_start=1
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 /protein_id="BA25600.1"
 /db_xref="GI:3059177"
 /translation="NAVESQGRPLVGLLCLVLPVVSNAKILLIPVDSHWSML
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 DSFLQRIKYKKIKDSAMLSGSHLLHNKELASLESFVMLDPLPCSPY
 AOVLSLPTVFPFLHALPCSLFEATQCPNPSYVPLSLSDPHMTFLQRYKNMLIAPS
 ONRICKVYSPTATLASFLQRETVQDLSASVWLFRSDPVMDYRPLPMNVFVG
 GINCLHNPESLQ"
 949..1190
 /partial
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 /note="does not fit consensus"
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gene
 TATA_signal
 exon
 gene
 CDS
 intron

Query Match 100.0%; Score 19; DB 91; Length 3341;
 Best Local Similarity 100.0%; Pred. No. 13;

Wed Jul 25 10:29:08 2001

us-09-142-095-4.rge

Page 8

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgctcccgccagaggtt 19
|||||
Db 3185 ttgctcccgccagaggtt 3167

Search completed: July 25, 2001, 05:17:10
Job time: 9233 sec

PT Improving drug trial efficiency comprises identifying participants

FT with Gilbert's syndrome - useful as their altered drug metabolism
 PT may hinder result interpretation

PS Claim 14, Page 13, 31pp: English.

CC This PCR primer (with primers AAT79542-44) flanks the TATA box sequence
 CC upstream of the uridine diphosphate glucuronosyltransferase (UGT) gene
 CC 1*1 exon 1 (see AAT79540), and was used to amplify fragments of 98-100
 CC bp. This gene is known to be associated with Gilbert's syndrome (GS). GS
 CC is a mild, common form of unconjugated hyperbilirubinaemia associated
 CC with reduced bilirubin glucuronidation capacity. Analysis of the genetic
 CC basis of GS has allowed 2 forms to be identified. One is a mild form
 CC associated with a homozygous 2 bp insertion in the TATA sequence
 CC upstream of the UGT*1 exon 1, and the other is a more severe form
 CC associated with heterozygosity for a mutation which, when homozygous,
 CC causes Crigler-Najjar type 2 disease. The first form is autosomal
 CC recessive and the second is inherited dominantly. Patients suffering from
 CC GS, which is benign, may have altered metabolism of some drugs, making it
 CC difficult to determine if an effect is due to the drug or the syndrome.
 CC Drug trial efficiency would be improved if potential participants can be
 CC screened for the genetic basis of GS, and eliminated or included on
 CC basis of them possessing or not possessing GS.

CC Sequence 19 BP; 2 A; 5 C; 5 G; 7 T; 0 other;

Query Match 100.0%; Score 19; DB 18; Length 19;

Best Local Similarity 100.0%; Pred. No. 1.5; Mismatches 0; Gaps 0;

OY 1 ttgtcctcgcagaggtt 19
 |||||
 DB 1 ttgtcctcgcagaggtt 19

RESULT 2

AAT79540/c
 ID AAT79540 standard; DNA; 620 BP.

XX AAT79540:

XX 23-JAN-1998 (first entry)

DE Upstream DNA sequence of UGT*1 gene exon 1.

XX uridine diphosphate glucuronosyltransferase gene; UGT;

XX Gilbert's syndrome; GS; unconjugated hyperbilirubinaemia;

XX bilirubin glucuronidation; Crigler-Najjar; type 2; drug metabolism;

XX Drug trial efficiency; screening; as.

XX Homo sapiens.

XX Key

XX protein_bind

XX GC_signal

XX misc-feature

XX protein_bind

XX protein_bind

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

XX misc-feature

FT protein_bind 386..392

FT /tag- g

FT /bound_moelty- AP1

FT /tag- h

FT /bound_moelty- AP1

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

FT protein_bind

FT /tag- j

FT /bound_moelty- HNF1

FT /tag- k

FT /note- "corresponds to positions -53 to -39 in patent"

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

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FT CDS

FT /tag- m

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FT protein_bind 386..392

FT /tag- g

FT /bound_moelty- AP1

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FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

FT protein_bind

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FT /tag- k

FT /note- "corresponds to positions -53 to -39 in patent"

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

FT TATA_signal

FT /tag- m

FT /codon_start- 612

FT CDS

FT /tag- m

FT /codon_start- 612

FT /tag- m

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FT /tag- m

FT protein_bind 386..392

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FT /bound_moelty- AP1

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

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FT /tag- j

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FT /note- "corresponds to positions -53 to -39 in patent"

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

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FT /tag- m

FT /codon_start- 612

FT CDS

FT /tag- m

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FT /codon_start- 612

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FT /tag- g

FT /bound_moelty- AP1

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FT explanation is given"

FT protein_bind

FT /tag- j

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FT explanation is given"

FT TATA_signal

FT /tag- m

FT /codon_start- 612

FT CDS

FT /tag- m

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FT /tag- m

FT protein_bind 386..392

FT /tag- g

FT /bound_moelty- AP1

FT /tag- h

FT /bound_moelty- AP1

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

FT protein_bind

FT /tag- j

FT /bound_moelty- HNF1

FT /tag- k

FT /note- "corresponds to positions -53 to -39 in patent"

FT /tag- 1

FT /note- "feature indicated in patent, but no further

FT explanation is given"

FT TATA_signal

FT /tag- m

FT /codon_start- 612

FT CDS

FT /tag- m

FT /codon_start- 612

FT /tag- m

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FT /codon_start- 612

FT /tag- m

FT /codon_start-

Db	605	TTCGCTCCTCCAGAGGTT	587
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ID	AAQ33024	standard; DNA; 1167 BP.	
XX	AAQ33024;		
XX	27-JAN-1993	(first entry)	
XX	UGT1A Exon 1 from the UGT1 gene locus.		
XX	DE		
XX	UGT1A; UGT1BP; UGTIC; UGTID; UGTLE; UGTIF; Isozyme; bilirubin;		
KW	UDP-glucuronosyl transferase; CN; SS.		
XX	Homo sapiens.		
OS			
XX	Key	Location/Qualifiers	
FH	misc_RNA	1..20	
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FT	/note-	"representation of 11.7 kbp of	
FT		non-sequenced DNA between the sequences	
FT		represented in AAQ33023 and AAQ33024"	
FT	TATA_signal	52..66	
FT	exon	/tag_b	
FT		166..909	
FT		/tag_c	
FT		/label-UGT1A_Exon_1	
FT	misc_RNA	181..909	
FT		/tag_d	
FT	/note-	"encodes transferase isoform; see CC"	
FT	misc_RNA	1148..1167	
FT		/tag_e	
FT		/note-	"representation of 5 kbp of
FT		non-sequenced DNA between the sequences	
FT		represented in AAQ33024 and AAQ33025"	
PN	WO9212987-A.		
XX	06-AUG-1992.		
PD			
XX	10-JAN-1992:	92WO-US00262.	
PX			
PR	10-JAN-1991:	91US-0639453.	
XX	(USSH) US DEPT HEALTH & HUMAN SERVICE.		
PA	Owens IS, Rittler JK;		
XI	WPI; 1992-284593/34.		
XX	P-PSDB; AAR30194.		
DR			
XX	Isolated gene locus UGT1, DNA segments and diagnostic probes -		
XX	types I and II		
PT	Disclosure; Fig 1F; 99p; English.		
PS			
XX	The isolated gene locus, UGT1, has a sequence of about 10000 bp		
CC	which represent (1) Exon 1, comprising 6 transcriptional units		
CC	(UGT1E, E, D, C, BP and A), represented in AAQ27366 and		
CC	AAQ33020-24 respectively;		
CC	(2) Exon 2, represented in AAQ33025;		
CC	(3) Exon 3, represented in AAQ33026;		
CC	(4) Exon 4, represented in AAQ33026;		
CC	(5) Exon 5, represented in AAQ33027; and		
CC	(6) about 69 kb of non-sequenced DNA.		
CC	Six unique N-terminal of 286-289 amino acids are encoded by		
CC	the six different first exons and identical C-terminal of 246 amino		
CC	acids are encoded by the common exons 2-5. The ugt1 gene locus		
CC	encodes a family of UDP-glucuronosyl transferase isozymes, two of		
CC	which metabolize bilirubin.		

CC Patients having Crigler-Najjar Syndrome (CN) Type I, have a
CC mutation present in the second common exon.
XX
SQ Sequence 1167 BP; 255 A; 259 C; 272 G; 340 T; 41 other:

OY 2 ttgcctcgtccagaggct 19
 |||||
DB 174 TTGCTCCTGCCAGACTT 157

RESULT 4
AAZ45058/c
ID AAZ45058 standard: DNA; 17 BP.
XX
AC AAZ45058;
XX
DT 28-FEB-2000 (first entry)
XX
DE Forward PCR primer used in the secondary amplification of UGT1 exon 1A.
XX
KM Uridine diphosphate-glucuronosyltransferase 1; UGT1; polymorphism; probe:
KM glucuronic acid; Crigler-Najjar syndrome; Gilbert syndrome; jaundice;
KM unconjugated hyperbilirubinemia; drug metabolism; transgenic animal;
KW pharmacogenetic screening; diagnosis; PCR primer; ss.
XX
OS Synthetic.
OS Homo sapiens.
XX
PN M09957322-A2.
XX
PD 11-NOV-1999.
XX
PF 04-MAY-1999; 99MO-US09702.
XX
PR 07-MAY-1998; 98US-0084807.
XX
PA (AAXS-) AXYS PHARM INC.
XX
PI Penny L, Galvin M;
XX
DR WPI: 2000-052981/04.
XX
PT New nucleic acid representing polymorphisms in the human uridine
PT diphosphate glucuronosyltransferase gene, used for diagnosis and
PT evaluation of drug metabolism -
XX
XX Examples: Page 16; 63pp; English.
XX
PS PCR primers AAZ45042-245073 are used to amplify human uridine
XX diphosphate-glucuronosyltransferase 1 (UGT1) exon sequences. The UGTs
CC are a family of enzymes that catalyse the glucuronic acid conjugation of
CC a wide range of endogenous and exogenous substrates including phenols,
CC alcohols, amines and fatty acids. Many of the reactions catalysed by
CC UGTs result in toxic substances being converted to compounds which are
CC more water soluble and are excreted. The invention relates to and
CC identifies UGT1 polymorphisms (AAZ45004-245041). The polymorphism
CC sequences are useful as probes for detecting UGT1 locus polymorphisms,
CC indicative of altered UGT1 expression or activity. These polymorphisms
CC are associated with Crigler-Najjar and Gilbert syndromes (unconjugated
CC hyperbilirubinemia) and drug metabolism. The genotyping of the UGT1 gene
CC is used to predict the rate of metabolism of UGT1 substrates, possible
CC drug-drug interactions and adverse side effects (i.e. to optimize drug
CC dosage), and to screen for diseases caused by exposure to toxins and to
CC study the effects of polymorphisms on enzymatic activity. The UGT1
CC sequences, including polymorphisms, can also be used to produce the
CC corresponding protein (or its fragments) or to generate transgenic
CC animals or modified cells e.g. for pharmacogenetic screening.

5Q Sequence 17 BP; 5 A; 4 C; 6 G; 2 T; 0 other;

Query Match 84.2%; Score 16; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgtctctgcagag 16
|||||
DB 16 TTGTCTCTGCCAGAG 1

RESULT 5
AAZ45074/c
ID AAZ45074 standard; DNA; 17 BP.

AC AAZ45074;

DT 28-FEB-2000 (first entry)

DE Forward PCR primer for sequencing UGT1 exon 1A polymorphism #1.

XX uridine diphosphate-glucuronosyltransferase 1; UGT1; polymorphism; probe;
KW glucuronic acid; Crigler-Najjar syndrome; Gilbert syndrome; jaundice;
KW unconjugated hyperbilirubinemia; drug metabolism; transgenic animal;
KW pharmacogenetic screening; diagnosis; PCR primer; ss.

XX Synthetic.

OS Homo sapiens.

PN MO9957322-A2.

PD 11-NOV-1999.

PF 04-MAY-1999; 99MO-US09702.

PR 07-MAY-1998; 98US-0084807.

PA (AXYS-) AXYS PHARM INC.

PI Penny L, Galvin M;

DR WPI: 2000-052981/04.

PT New nucleic acid representing polymorphisms in the human uridine
diphosphate glucuronosyltransferase gene, used for diagnosis and
evaluation of drug metabolism -

PS Examples: Page 19; 63pp; English.

CC Primers AAZ45074-245109 are used to sequence the human uridine
diphosphate-glucuronosyltransferase 1 (UGT1) exon polymorphism
sequences. The UGTs are a family of enzymes that catalyze the glucuronic
acid conjugation of a wide range of endogenous and exogenous substrates
including phenols, alcohols, amines and fatty acids. Many of the
reactions catalyzed by UGTs result in toxic substances being converted
to compounds which are more water soluble and are excreted. The
invention relates to and identifies UGT1 polymorphisms (AAZ45004-245041).
CC The polymorphism sequences are useful as probes for detecting UGT1 locus
polymorphisms, indicative of altered UGT1 expression or activity. These
polymorphisms are associated with Crigler-Najjar and Gilbert syndromes
(unconjugated hyperbilirubinemia) and drug metabolism. The genotyping
of the UGT1 gene is used to predict the rate of metabolism of UGT1
substrates, possible drug-drug interactions and adverse side effects
(i.e. to optimize drug dosage), and to screen for diseases caused by
exposure to toxins and to study the effects of polymorphisms on
enzymatic activity. The UGT1 sequences, including polymorphisms on
be used to produce the corresponding protein (or its fragments) or to
generate transgenic animals or modified cells e.g. for pharmacogenetic
screening.

XX Sequence 17 BP; 5 A; 4 C; 6 G; 2 T; 0 other;

Query Match 84.2%; Score 16; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ttgtctctgcagag 16
|||||
DB 16 TTGTCTCTGCCAGAG 1

RESULT 6
AAZ45200/c
ID AAZ45200 standard; DNA; 1307 BP.

AC AAZ45200;

DT 15-MAY-2001 (first entry)

DE Human 2hdelcalp DNA.

XX Notch receptor; ligand; cancer; melanoma; ischemia; ds.

XX Homo sapiens.

PN WO200112664-A2.

PD 22-FEB-2001.

PF 17-AUG-2000; 2000MO-US22609.

PR 19-AUG-1999; 99US-0149934.

PA (CHIR) CHIRON CORP.

PI Vivien C, Rohan M, Williams LT;

DR WPI: 2001-211201/21.

PT Novel notch receptor ligands useful for modulating angiogenesis and
immune responses for treating rheumatoid arthritis, cancer-related
angiogenesis to stop tumor growth, and as diagnostic reagents -

PS Disclosure: Fig 6; 66pp; English.

CC The present invention relates to a novel Notch receptor ligand.
CC The invention is useful for detecting Notch ligand expression in
human cancer cells or melanoma cells. Also useful for enhancing
angiogenesis in a mammal, useful when the mammal exhibits
tissue ischemia.

SO Sequence 1307 BP; 294 A; 328 C; 396 G; 289 T; 0 other;

Query Match 84.2%; Score 16; DB 22; Length 1307;
Best Local Similarity 100.0%; Pred. No. 75;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 gctctgcagagct 19
|||||
DB 1119 GCTCTGCCAGAGCTT 1104

RESULT 7

ID AAZ4518/c

AC AAZ4518;

DT 12-OCT-1999 (first entry)

DE Human gene expression product cDNA sequence SEQ ID NO: 3988.

XX Human; gene; gene expression product; diagnosis; therapy; probe;

XX detection; mapping; tissue typing; profiling; forensic; cancer;
 KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.
 OS Homo sapiens.
 XX MO9938972-A2.
 XX 05-AUG-1999.
 XX 28-JAN-1999; 99MO-US01619.
 XX 03-APR-1998; 98US-0080666.
 XX 28-JAN-1998; 98US-0072910.
 XX 24-FEB-1998; 98US-0075954.
 XX 31-MAR-1998; 98US-0080114.
 XX 03-APR-1998; 98US-0080515.
 PA (CHIR) CHIRON CORP.
 PA (HYSE-) HYSEQ INC.
 XX Ctkvenjakov R, Dickson M, Dermanac R, Dermanac S;
 PI Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;
 PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;
 PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
 PI Stache-Crain B, Sudduth-Klinger J, Williams LP;
 XX WPI; 1999-494092/41.
 DR Novel human genes and their expression products which are
 XX differentially expressed in different cell types
 PT Claim 1; Page 1892-1893; 2479pp; English.
 XX The present invention describes a library of human polynucleotides
 CC comprising the sequences given in AA21532 to AA21779. Also described is
 CC a method of detecting differentially expressed genes correlated with the
 CC cancerous state of a mammalian cell, comprising detecting at least one
 CC differentially expressed gene product in a test sample from a cell
 CC suspected of being cancerous, where the gene product is encoded by one
 CC of the 5248 polynucleotide sequences given in AA21532 to AA21779. The
 CC polynucleotides can be used as a source of primers and probes, which can
 CC be used for a variety of purpose, e.g. detection of expression levels,
 CC mapping, tissue typing or profiling, forensics, genetic analysis and
 CC detection of polymorphisms. Polypeptides encoded by the polynucleotides
 CC can be used for raising antibodies for experimental, diagnostic and
 CC therapeutic purposes. The polynucleotides may also be used to construct
 CC arrays for diagnostics (which may be used to determine function of an
 CC encoded protein); and to detect differences in expression levels between
 CC two cells (e.g. to identify abnormal or diseased tissue in a human, to
 CC identify a genetic predisposition or susceptibility to a disease such as
 CC cancer). The polynucleotides of the invention are especially used in the
 CC diagnosis, prognosis and management of colorectal cancer, breast cancer,
 CC and lung cancer. The polynucleotides can also be used to screen for
 CC peptide analogues and antagonists.
 XX Sequence 457 BP; 120 A; 88 C; 93 G; 139 T; 17 other;
 SO

Query Match 83.2%; Score 15.8; DB 20; Length 457;
 Best Local Similarity 89.5%; Pred. No. 83;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
 |||||
 DB 300 TTGTCTCTGCGAGAGCTT 282

RESULT 8
 AAV22955/c
 ID AAV22955 standard; CDNA; 843 BP.
 XX AAV22955;
 XX

DT 04-AUG-1998 (first entry)
 XX CDNA encoding human bone morphogenetic protein-16 (BMP-16).
 DE Human; bone morphogenetic protein-16; BMP-16; murine protein; nodal;
 KW formation; bone; cartilage; treatment; wound healing; reduction;
 KW fibrosis; scar tissue formation; ss.
 XX Homo sapiens.
 OS
 XX Key Location/Qualifiers
 XX CDS 1..843
 XX sig_peptide 1..510
 XX mat_peptide 511..840
 XX /tag- b
 XX /tag- c
 XX MO9812322-A1.
 XX 26-MAR-1998.
 XX 09-JUL-1997; 97MO-US11954.
 XX 18-SEP-1996; 96US-0715202.
 XX (GENY) GENETICS INST INC.
 XX Celeste AJ, Murray BL;
 PI WPI; 1998-217262/19.
 DR P-PSDB; AAW56477.
 XX New isolated bone morphogenetic protein-16 - used to develop
 PT products for inducing formation of bone, cartilage and other
 PT connective tissue, particularly for wound healing and tissue repair
 XX Claim 5; Pages 33-34; 43pp; English.
 XX The present sequence encodes a human bone morphogenetic protein-16
 CC (BMP-16). Human BMP-16 is a homologue of a murine protein called nodal,
 CC which is expressed in the mouse node during gastrulation. BMP-16 cDNA is
 CC isolated from a human genomic library screened with a probe derived from
 CC the nodal cDNA sequence. The BMP-16 proteins can induce the formation of
 CC bone, cartilage or other connective tissue. They can be used for treating
 CC bone, cartilage or other connective tissue defects, periodontal disease
 CC or healing of various types of tissues and wounds. They can also increase
 CC neuronal, astrocytic and glial cell survival and therefore be useful in
 CC transplantation and treatment of conditions exhibiting a decrease in
 CC neuronal survival and repair. They can also exhibit properties such as
 CC angiogenic, chemotactic and/or chemoattractant properties, and effects
 CC on cells including induction of collagen synthesis, fibrosis, and effects
 CC on differentiation responses, cell proliferative responses and responses
 CC involving cell adhesion, migration and extracellular matrices. These
 CC properties make the proteins potential agents for wound healing,
 CC reduction of fibrosis and reduction of scar tissue formation.
 XX Sequence 843 BP; 176 A; 241 C; 244 G; 182 T; 0 other;
 SO

Query Match 83.2%; Score 15.8; DB 19; Length 843;
 Best Local Similarity 89.5%; Pred. No. 89;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
 |||||
 DB 412 TTGTCTCTGCGAGAGCTT 394

RESULT 9
 AAV6866/c
 ID AAV6866 standard; DNA; 1095 BP.
 XX AAV6866;
 XX

```

AC      AA68666;
XX
DT      03-JUN-1999      (first entry)
XX
DE      Nucleotide sequence of the human Tango-78 gene.
XX
XX      Human; Tango-78; host cell; recombinant protein; antibody;
XX      receptor; specific binding agent; probe; primer; hybridisation;
XX      amplification; mutation; genetic mapping; ss.
XX
OS      Homo sapiens.
XX
XX      Key      Location/Qualifiers
XX      FT      CDS      588..1094
XX      FT      /tag=a
XX      FT      /product="Tango-78"
XX
XX      M09306427-A1.
XX
XX      11-FEB-1999.
XX
XX      04-AUG-1998;      98MO-US1241.
XX
XX      04-AUG-1997;      97US-0054645.
XX
XX      (MILL-) MILLENNIUM BIOTHERAPEUTICS INC.
XX
XX      McCarthy SA;
XX
XX      WPI; 1999-153693/13.
XX      P-PSDB; AAW84595.
XX
XX      New nucleic acid encoding human Tango-78, -79 and -81 proteins -
XX      useful for diagnosis and treatment of Tango-associated diseases
XX
XX      Claim 1; Fig 1; 67pp; English.
XX
XX      This is the nucleotide sequence encoding the human Tango-78
XX      protein used in the method of the invention. Host cells containing
XX      the Tango protein are used to produce recombinant proteins for
XX      raising antibodies. It is also used in identifying specific
XX      binding agents (including cognate receptors), which can be used to
XX      determine amounts of recombinant protein in cells or
XX      therapeutically. Antibodies or other specific binding agents, are
XX      used to detect recombinant proteins and fragments of the Tango
XX      nucleotide sequence can be used as probes or primers for detecting
XX      the Tango gene, specifically mRNA, in usual hybridisation or
XX      amplification assays. These assays are used for diagnosis of
XX      diseases associated with abnormal expression of Tango proteins,
XX      e.g. detecting mutations in the Tango gene. Fragments of the Tango
XX      nucleic acid sequence are also used for genetic mapping and
XX      chromosome identification, and as antisense, ribozyme or
XX      triplex-forming therapeutics. Antibodies may also be used to generate
XX      anti-idiotypic antibodies.
XX
XX      Sequence 1095 BP; 231 A; 296 C; 312 G; 256 T; 0 other;
XX
XX
XX      Query Match      83.2%; Score 15.8; DB 20; Length 1095;
XX      Best Local Similarity 89.5%; Pred. No. 92;
XX      Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX      1 ttgtgctctcgagaggtt 19
XX      | ||||| |||||
XX      Db      801 tctgctctcgcgaggtt 793
XX
XX      RESULT 10
XX      ID      AAX31924/c
XX      XX      AAX31924 standard; DNA; 1156 BP.
XX
XX      AC      AAX31924;
XX

```

DT		18-JUN-1999	(first entry)
XX			
DE			Human nodal protein encoding DNA.
KW	Nodal protein; lefty protein; TGF-beta; sexual development; human; bone;		
KM	pituitary; cartilage; osteoarthritis; osteoporosis; haematopoiesis;		
KW	interstitial lung disease; wound healing; tissue repair; cancer;		
KX	immunosuppression; inflammatory bowel disease; lymphoma; immunity;		
KM	infectious disease; ss.		
OS	Homo sapiens.		
XX			
FH	Key	Location/Qualifiers	
FT	CDS	1..852	
FT	/ftag-	a	
FT	/product=	"Nodal protein"	
FT	/note=	"the start codon is not indicated"	
FT	misc-feature	517..849	
FT	/ftag-	b	
FT	/note=	"sequence coding for the active fragment of the Nodal polypeptide"	
PX	WP09909198-A1.		
PD	25-FEB-1999.		
PF	20-AUG-1998:	98WO-US17211.	
PR	21-AUG-1997:	9TUS-0056565.	
PA	(HOMA-) HUMAN GENOME SCI INC.		
PI	Ebner R, Ruben SM, Soppet DR;		
DR	Mpi: 1999-190173/L16.		
DX	P-PDB: AAI03849.		
PT	New isolate human Nodal and Lefty polypeptides		
PS	Claim 2; Fig 1A; 182pp; English.		
CC	The present invention relates to novel human nodal and lefty proteins which are members of the TGF-beta family. The human nodal and lefty proteins may be involved in a developmental process such as the correct formation of various structures or in one or more post-developmental capacities including sexual development, pituitary hormone production, and the creation of bone and cartilage. The nodal and lefty polypeptides are useful for enhancing or enriching the growth and/or differentiation of specific cell populations, eg. embryonic cells or stem cells. They can be used to treat such conditions as osteoarthritis, osteoporosis, and other abnormalities of bone, cartilage, muscle, tendon, ligament, and/or other connective tissues and/or organs such as liver, lung, cardiac, pancreas, and kidney. Compositions containing nodal and lefty proteins may be useful for growth promotion, for treating periodontal disease and for modulating hematopoiesis, wound healing and tissue repair. They can also be used for the treatment of tumours, cancers, interstitial lung disease, and any dysregulation of the growth and differentiation patterns of cell function including autoimmunity, arthritis, leukaemia, lymphomas, immunosuppression, immunity, humoral immunity, inflammatory bowel disease, myelosuppression, or infectious diseases. The present sequence represents a DNA encoding a human nodal polypeptide. The cDNA encoding the nodal protein is deposited under the ATCC deposit No. 209092 and/or 209135.		
SQ	Sequence 1156 BP; 285 A; 298 C; 340 G; 233 T; 0 Other;		
Query Match	83.2%; Score 15.8; DB 20; Length 1156;		
Best Local Similarity	89.5%; Pied NO. 93;		
Matches 17; Conservative	0; Mismatches 2; Indels 0; Gaps 0		
y	1 ttgtctcgcgccagaagt t		

|||||
421 TCTGCTCTCCAGAGGTT 403

RESULT 11

AA27725/C
ID: AAF1725 standard; CDNA: 1571 BP.

XX AAF27725;

XX 28-MAR-2001 (first entry)

XX Human transport protein TPT-25 coding sequence.

XX Human: transport protein; TPT: transport disorder; metabolic disorder;

XX neurological disorder; cardiovascular disorder; reproductive disorder;

XX immune disorder; cancer; ss.

XX Homo sapiens.

XX MO200078953-r2.

XX 28-DEC-2000.

XX 16-JUN-2000; 2000MO-US16668.

XX 17-JUN-1999; 99US-013923.

XX 10-AUG-1999; 99US-014817.

XX 18-AUG-1999; 99US-0149357.

XX 28-OCT-1999; 99US-0162287.

XX (INCYTE GENOMICS INC.

XX La1 P, Yang J, Yue H, Hillman JL, Tang YT, Sandman G, Burford N;

XX Baughn MR, Alexander Y, Lu RM, Au-Yang J, Patterson C;

XX WPI: 2001-041424/05.

XX P-PSDB: AAB60105.

XX Isolated polypeptide with a human transport protein sequence is useful

XX for the diagnosis, prevention and treatment of disorders associated

XX with the immune, reproductive, and cardiovascular systems.

XX Claim 5: Page 155-156; 16pp; English.

XX The present invention provides the protein and coding sequences for 43

XX novel human transport proteins (designated TPTs). These can be used in

XX the diagnosis and treatment of transport, metabolic, neurological,

XX reproductive, cardiovascular, and cell proliferative

XX disorders such as cancer.

XX Sequence 1571 BP; 572 A; 265 C; 289 G; 495 T; 0 other:

XX Query Match 83.2% Score 15.8; DB 22; Length 1571;

XX Best Local Similarity 89.5% Pred. No. 96;

XX Matches 17; Cons: 17; Mismatches 2; Indels 0; Gaps 0;

XX DB 1571 TCTGCTCTCCAGAGGTT 1576

RESULT 12

AA27725/C
ID: AAC98907 standard; CDNA: 1945 BP.

XX AAC98907;

XX 09-MAR-2001 (first entry)

XX Human: pancreatic cancer antigen nucleotide sequence SEQ ID NO:135.

XX 04-AUG-1998 (first entry)

XX Human: pancreas; pancreatic cancer; pancreatic cancer antigen;

XX detection; diagnosis; identification; cytostatic; neuroprotective;

XX neoplastic; immunomodulatory; relaxant; contraceptive; gynaecological;

XX anti-inflammatory; cardiant; gene therapy; chromosome mapping;

XX linkage analysis; tissue identification; tissue typing; forensic;

XX neutral; immune system; muscular; reproductive; gastrointestinal;

XX pulmonary; cardiovascular; renal; proliferative; ss.

XX Homo sapiens.

XX MO200055320-AL.

XX 21-SEP-2000.

XX 08-MAR-2000; 2000MO-US05989.

XX 12-MAR-1999; 99US-0124270.

XX (HUMA-) HUMAN GENE SCI INC.

XX Rosen CA, Ruben SM;

XX WPI: 2000-579444/54.

XX P-PSDB: AAB54142.

XX New nucleic acid that is a pancreatic cancer antigen for preventing,

XX treating, or ameliorating a medical condition, particular pancreatic

XX cancer, or for use in assays for diagnosing a pathological condition.

XX Claim 1: Page 594-595; 1379pp; English.

XX AAC98773 to AAC99231 encode the human pancreatic cancer associated

XX proteins, called pancreatic cancer antigens, given in AAB54008 to

XX AAB54466. The human pancreatic cancer antigens have cytostatic,

XX neuroprotective, neoplastic, immunomodulatory, relaxant, contraceptive,

XX gynaecological, cardiant and anti-inflammatory activities, and can be used

XX in gene therapy. The polynucleotide and proteins can be used for

XX preventing, treating, or ameliorating a medical condition or in assays

XX for diagnosing a pathological condition or a susceptibility to one in a

XX subject. Binding partners to the proteins and the activity of the

XX proteins can be identified. The pancreatic cancer antigens can be used to

XX detect, treat or prevent pancreatic disorders, especially cancer.

XX Agonists and antagonists to the antigens can be screened for. The

XX pancreatic cancer antigen polynucleotide can be used to design nucleic

XX acid hybridization probes that can be used in chromosome mapping, linkage

XX analysis, tissue identification and/or typing and a variety of forensic

XX and diagnostic methods. The proteins can be used to generate antibodies

XX which are used to purify, detect and target the polypeptides, including

XX both in vivo and in vitro for diagnostic and therapeutic methods. The

XX proteins can be used to treat, prevent, neutral, immune system, muscular,

XX reproductive, gastrointestinal, pulmonary, cardiovascular, renal or

XX proliferative disorders. AAC99232 to AAC99240 and AAB54467 represent

XX sequences used in the exemplification of the present invention.

XX Sequence 1945 BP; 533 A; 300 C; 477 G; 552 T; 13 other:

XX Query Match 83.2% Score 15.8; DB 21; Length 1945;

XX Best Local Similarity 89.5% Pred. No. 98;

XX Matches 17; Cons: 17; Mismatches 2; Indels 0; Gaps 0;

XX DB 1203 TTGCTCTCCAGAGGTT 1185

RESULT 13

AA22956/C
ID: AAV22956 standard; DNA: 2003 BP.

XX AAV22956;

XX 04-AUG-1998 (first entry)

PR 29-MAR-1999; 99US-0126785.
PR 01-APR-1999; 99US-0127462.
PR 06-APR-1999; 99US-0128234.
PR 08-APR-1999; 99US-0128714.
PR 16-APR-1999; 99US-0129845.
PR 19-APR-1999; 99US-0130077.
PR 21-APR-1999; 99US-0130449.
PR 23-APR-1999; 99US-0130510.
PR 23-APR-1999; 99US-0130891.
PR 28-APR-1999; 99US-0131449.
PR 30-APR-1999; 99US-0132048.
PR 04-MAY-1999; 99US-0132407.
PR 05-MAY-1999; 99US-0132484.
PR 06-MAY-1999; 99US-0132485.
PR 06-MAY-1999; 99US-0132486.
PR 07-MAY-1999; 99US-0132487.
PR 11-MAY-1999; 99US-0134286.
PR 14-MAY-1999; 99US-0134218.
PR 14-MAY-1999; 99US-0134219.
PR 14-MAY-1999; 99US-0134221.
PR 14-MAY-1999; 99US-0134370.
PR 18-MAY-1999; 99US-0134768.
PR 19-MAY-1999; 99US-0134941.
PR 20-MAY-1999; 99US-0135124.
PR 21-MAY-1999; 99US-0135353.
PR 24-MAY-1999; 99US-0135629.
PR 25-MAY-1999; 99US-0136021.
PR 27-MAY-1999; 99US-0136392.
PR 28-MAY-1999; 99US-0136782.
PR 01-JUN-1999; 99US-0137222.
PR 03-JUN-1999; 99US-0137528.
PR 04-JUN-1999; 99US-0137502.
PR 07-JUN-1999; 99US-0137724.
PR 08-JUN-1999; 99US-0138094.
PR 10-JUN-1999; 99US-0138540.
PR 10-JUN-1999; 99US-0138647.
PR 14-JUN-1999; 99US-0139119.
PR 16-JUN-1999; 99US-0139452.
PR 17-JUN-1999; 99US-0139453.
PR 18-JUN-1999; 99US-0139454.
PR 18-JUN-1999; 99US-0139455.
PR 18-JUN-1999; 99US-0139456.
PR 18-JUN-1999; 99US-0139457.
PR 18-JUN-1999; 99US-0139458.
PR 18-JUN-1999; 99US-0139459.
PR 18-JUN-1999; 99US-0139460.
PR 18-JUN-1999; 99US-0139461.
PR 18-JUN-1999; 99US-0139462.
PR 18-JUN-1999; 99US-0139463.
PR 18-JUN-1999; 99US-0139750.
PR 18-JUN-1999; 99US-0139763.
PR 21-JUN-1999; 99US-0139817.
PR 22-JUN-1999; 99US-0139899.
PR 23-JUN-1999; 99US-0140353.
PR 23-JUN-1999; 99US-0140354.
PR 24-JUN-1999; 99US-0140354.
PR 26-JUN-1999; 99US-0140695.
PR 29-JUN-1999; 99US-0140823.
PR 30-JUN-1999; 99US-0140991.
PR 01-JUL-1999; 99US-0141287.
PR 01-JUL-1999; 99US-0141842.
PR 02-JUL-1999; 99US-0142154.
PR 06-JUL-1999; 99US-0142055.
PR 08-JUL-1999; 99US-0142290.
PR 09-JUL-1999; 99US-0142803.
PR 12-JUL-1999; 99US-0142920.
PR 13-JUL-1999; 99US-0143977.
PR 14-JUL-1999; 99US-0143542.
PR 15-JUL-1999; 99US-0143624.
PR 16-JUL-1999; 99US-0144003.
PR 16-JUL-1999; 99US-0144083.
PR 16-JUL-1999; 99US-0144086.

PR 19-JUL-1999; 99US-0144325.
PR 19-JUL-1999; 99US-0144331.
PR 19-JUL-1999; 99US-0144332.
PR 19-JUL-1999; 99US-0144333.
PR 19-JUL-1999; 99US-0144334.
PR 19-JUL-1999; 99US-0144335.
PR 20-JUL-1999; 99US-0144352.
PR 20-JUL-1999; 99US-0144632.
PR 21-JUL-1999; 99US-0144884.
PR 21-JUL-1999; 99US-0144814.
PR 21-JUL-1999; 99US-0145086.
PR 22-JUL-1999; 99US-0145086.
PR 22-JUL-1999; 99US-0145087.
PR 22-JUL-1999; 99US-0145089.
PR 23-JUL-1999; 99US-0145192.
PR 23-JUL-1999; 99US-0145145.
PR 23-JUL-1999; 99US-0145218.
PR 23-JUL-1999; 99US-0145224.
PR 26-JUL-1999; 99US-0145276.
PR 27-JUL-1999; 99US-0145913.
PR 27-JUL-1999; 99US-0145918.
PR 28-JUL-1999; 99US-0145919.
PR 02-AUG-1999; 99US-0146386.
PR 02-AUG-1999; 99US-0146388.
PR 03-AUG-1999; 99US-0147038.
PR 04-AUG-1999; 99US-0147204.
PR 05-AUG-1999; 99US-0147302.
PR 05-AUG-1999; 99US-0147192.
PR 05-AUG-1999; 99US-0147260.
PR 06-AUG-1999; 99US-0147303.
PR 06-AUG-1999; 99US-0147416.
PR 09-AUG-1999; 99US-0147493.
PR 09-AUG-1999; 99US-0147935.
PR 10-AUG-1999; 99US-0148171.
PR 11-AUG-1999; 99US-0148319.
PR 12-AUG-1999; 99US-0148341.
PR 13-AUG-1999; 99US-0148684.
PR 16-AUG-1999; 99US-0148368.
PR 17-AUG-1999; 99US-0148175.
PR 18-AUG-1999; 99US-0148426.
PR 20-AUG-1999; 99US-0148722.
PR 20-AUG-1999; 99US-0148723.
PR 20-AUG-1999; 99US-0148929.
PR 23-AUG-1999; 99US-0148902.
PR 25-AUG-1999; 99US-0150566.
PR 26-AUG-1999; 99US-0150588.
PR 27-AUG-1999; 99US-0151065.
PR 27-AUG-1999; 99US-0151066.
PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151436.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
PR 10-SEP-1999; 99US-0153070.
PR 13-SEP-1999; 99US-0153758.
PR 15-SEP-1999; 99US-0154018.
PR 16-SEP-1999; 99US-0154038.
PR 20-SEP-1999; 99US-0154779.
PR 22-SEP-1999; 99US-0155139.
PR 23-SEP-1999; 99US-0155486.
PR 24-SEP-1999; 99US-0155659.
PR 28-SEP-1999; 99US-0156458.
PR 29-SEP-1999; 99US-0156596.
PR 04-OCT-1999; 99US-0157117.
PR 05-OCT-1999; 99US-0157753.
PR 06-OCT-1999; 99US-0157866.
PR 07-OCT-1999; 99US-0158029.
PR 08-OCT-1999; 99US-0158233.

XX	Key	Location/Qualifiers
FT	CD5	244..2628
FT		/*tag- a
XX	CA2139384-A.	
XX	12-NOV-1995.	
XX	30-DEC-1994;	94CA-2139384.
XX	11-MAY-1994;	94GB-0009396.
XX	(CIBA) CIBA GEIGY AG.	
XX	Gouilleux F, Groner B, Makao H;	
XX	WPI, 1996-077786/09.	
XX	P-PADB; AAR88199.	
XX	DNA encoding mammary gland factor protein - used to identify cpds.	
XX	affecting intracellular signal transduction of a lactogenic hormone,	
XX	or cytokine	
XX	Claim 2, Page 28-34; 42pp; English.	
XX	A cDNA clone (AAT10554) coding for mammary gland factor (MGF)	
XX	(AAR88199) was obt'd. from a cDNA library derived from sheep	
XX	lactating mammary tissue mRNA using probes (AAT10555-56) based on	
XX	internal peptides of MGF. The cDNA is used to produce recombinant	
XX	MGF in host cells, or as a probe. Transgenic animals, e.g. sheep,	
XX	overexpressing MGF can be produced in order to increase milk prodn.	
XX	or to produce a therapeutically useful protein.	
XX	Sequence 2818 BP; 618 A; 861 C; 819 G; 520 T; 0 other.	
XX	Query Match	83.2%; Score 15.8; DB 17; Length 2818;
XX	Best Local Similarity	89.5%; Pred. No. 1e+02;
XX	Matches 17; Conservative	0; Mismatches 2; Indels 0; Gaps 0;
XX	1 ttgtctctctgcagaggtt 19	
XX		
XX	2599 ttattctctgcagagcgt 2617	
XX	RESULT 15	
XX	AAC47622	
XX	ID AAC47622 standard; DNA; 3346 BP.	
XX	AAC47622;	
XX	18-OCT-2000 (first entry)	
XX	Arabidopsis thaliana DNA fragment SEQ ID NO: 54504.	
XX	Hybridisation assay; genetic mapping; gene expression control;	
XX	protein identification; signal transduction pathway;	
XX	metabolic pathway; promoter; termination sequence; ss.	
XX	Arabidopsis thaliana.	
XX	EP1033405-A2.	
XX	06-SEP-2000.	
XX	25-FEB-2000; 2000EP-0301439.	
XX	25-FEB-1999;	99US-0121825.
XX	05-MAR-1999;	99US-0123180.
XX	09-MAR-1999;	99US-0123548.
XX	23-MAR-1999;	99US-0125788.
XX	25-MAR-1999;	99US-0126264.

PR 12-OCT-1999: 99US-0158369.
PR 13-OCT-1999: 99US-0159293.
PR 13-OCT-1999: 99US-0159294.
PR 13-OCT-1999: 99US-0159295.
PR 14-OCT-1999: 99US-0159329.
PR 14-OCT-1999: 99US-0159330.
PR 14-OCT-1999: 99US-0159331.
PR 14-OCT-1999: 99US-0159637.
PR 14-OCT-1999: 99US-0159638.
PR 18-OCT-1999: 99US-0159584.
PR 21-OCT-1999: 99US-0160741.
PR 21-OCT-1999: 99US-0160767.
PR 21-OCT-1999: 99US-0160768.
PR 21-OCT-1999: 99US-0160770.
PR 21-OCT-1999: 99US-0160814.
PR 21-OCT-1999: 99US-0160815.
PR 22-OCT-1999: 99US-0160980.
PR 22-OCT-1999: 99US-0160981.
PR 22-OCT-1999: 99US-0160989.
PR 25-OCT-1999: 99US-0161404.
PR 25-OCT-1999: 99US-0161405.
PR 25-OCT-1999: 99US-0161406.
PR 26-OCT-1999: 99US-0161359.
PR 26-OCT-1999: 99US-0161360.
PR 26-OCT-1999: 99US-0161361.
PR 28-OCT-1999: 99US-0161920.
PR 28-OCT-1999: 99US-0161992.
PR 28-OCT-1999: 99US-0161993.
PR 29-OCT-1999: 99US-0162142.

Query Match 83.2%; Score 15.8; DB 21; Length 3346;
Best Local Similarity 89.5%; Pred. No. 1e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggtt 19
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Db 1742 ttgtctctgcagaggtt 1760

Search completed: July 25, 2001, 05:23:06
Job time: 4679 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:18:53 / Search time 117.39 seconds
(without alignments)
29.985 Million cell updates/sec

Title: US-09-142-095-4

Sequence: 1 ttgtctctgcagaggtt 19

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 317530 seqs, 92630169 residues

Total number of hits satisfying chosen parameters: 635060

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

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2: /cgn2_6/ptodata/2/1na/5b_COMB.seq:*
3: /cgn2_6/ptodata/2/1na/6a_COMB.seq:*
4: /cgn2_6/ptodata/2/1na/6b_COMB.seq:*
5: /cgn2_6/ptodata/2/1na/PCTUS_COMB.seq:*
6: /cgn2_6/ptodata/2/1na/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	19	100.0	1190	5	PCT-US92-00282-18
C 2	15.8	83.2	843	2	US-08-715-202A-1
C 3	15.8	83.2	2002	2	US-08-715-202A-3
C 4	15.8	83.2	2818	1	US-08-366-276-1
C 5	14.8	77.9	2282	1	US-08-373-579-5
C 6	14.8	77.9	2282	1	US-08-418-595-5
C 7	14.8	77.9	2282	1	US-08-665-926-5
C 8	14.8	77.9	2282	4	US-09-162-437-5
C 9	14.8	77.9	5948	2	US-08-447-411-44
C 10	14.8	77.9	5948	2	US-08-662-227-1
C 11	14.4	75.8	1858	2	US-08-359-705B-7
C 12	14.4	75.8	1858	2	US-08-286-846A-7
C 13	14.4	75.8	1858	2	US-08-457-880A-7
C 14	14.4	75.8	1858	3	US-08-444-622A-7
C 15	14.4	75.8	1858	3	US-08-942-562-7
C 16	14.4	75.8	1858	4	US-09-156-923-7
C 17	14.4	75.8	2715	2	US-08-359-705B-5
C 18	14.4	75.8	2715	2	US-08-286-846A-5
C 19	14.4	75.8	2715	2	US-08-457-880A-5
C 20	14.4	75.8	2715	3	US-08-444-622A-5
C 21	14.4	75.8	2715	3	US-08-942-562-5
C 22	14.4	75.8	2715	4	US-09-156-923-5
C 23	14.4	75.8	2940	2	US-08-286-846A-8
C 24	14.4	75.8	2940	2	US-08-441-104A-8
C 25	14.4	75.8	2940	2	US-08-440-816A-8
C 26	14.2	74.7	300	2	US-07-938-154-6
C 27	14.2	74.7	300	5	PCT-US91-03311-6

28	14.2	74.7	693	1	US-08-052-205-10	Sequence 10, Appl
29	14.2	74.7	693	1	US-08-595-974-10	Sequence 10, Appl
30	14.2	74.7	759	1	US-08-052-205-8	Sequence 8, Appl1
31	14.2	74.7	759	1	US-08-595-974-8	Sequence 8, Appl1
32	14.2	74.7	1044	1	US-08-052-205-6	Sequence 6, Appl1
33	14.2	74.7	1044	1	US-08-595-974-6	Sequence 6, Appl1
34	14.2	74.7	1062	1	US-08-052-205-1	Sequence 1, Appl1
35	14.2	74.7	1062	1	US-08-595-974-1	Sequence 1, Appl1
36	14.2	74.7	1110	1	US-08-052-205-5	Sequence 5, Appl1
37	14.2	74.7	1110	1	US-08-595-974-5	Sequence 5, Appl1
38	14.2	74.7	1393	1	US-08-052-205-2	Sequence 2, Appl1
39	14.2	74.7	1393	1	US-08-595-974-2	Sequence 2, Appl1
40	14.2	74.7	1449	2	US-08-705-868-2	Sequence 2, Appl1
41	14.2	74.7	1449	2	US-09-123-635-2	Sequence 2, Appl1
42	14.2	74.7	1451	1	US-08-031-143B-68	Sequence 66, Appl
43	14.2	74.7	1470	1	US-08-052-205-3	Sequence 3, Appl1
44	14.2	74.7	1470	1	US-08-595-974-3	Sequence 3, Appl1
45	14.2	74.7	1608	2	US-08-424-224-1	Sequence 1, Appl1

ALIGNMENTS

RESULT 1
PCT-US92-00282-18/c

Sequence 18, Application PC/TUS9200282

GENERAL INFORMATION:

APPLICANT: OMENS, IDA S.

APPLICANT: RITTER, JOSEPH K.

TITLE OF INVENTION: THE GENETIC LOCUS UGT1 AND A MUTATION

TITLE OF INVENTION: THEREN.

NUMBER OF SEQUENCES: 40

CORRESPONDENCE ADDRESS:

ADDRESSEE: CUSHMAN DABRY & CUSHMAN

STREET: 1615 L STREET, N.W.

CITY: WASHINGTON

STATE: D.C.

COUNTRY: U.S.A.

ZIP: 20036-5601

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: PCT/US92/00282

FILING DATE: 19920110

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: SCOTT, WATSON T.

REGISTRATION NUMBER: 26581

REFERENCE/DOCKET NUMBER: 91532-PCT

TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-861-3000

TELEFAX: 202-822-0944

TELEX: 6714627 CUSH

INFORMATION FOR SEQ. ID NO. 18:

SEQUENCE CHARACTERISTICS:

LENGTH: 1190 base pairs

TYPE: NUCLEIC ACID

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: CDNA

PCT-US92-00282-18

Query Match 100.0%; Score 19; DB 5; Length 1190;

Best Local Similarity 100.0%; Pred. No. 0.49;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ttgtctctgcagaggtt 19

78 ttgtctctgcagaggtt 60

RESULT 2
US-08-715-202A-1/c
Sequence 1, Application US/08715202A
Patent No. 5363403
GENERAL INFORMATION:
APPLICANT: CELESTE, ANTHONY J.
APPLICANT: MURRAY, BETH L.
TITLE OF INVENTION: BONE MORPHOGENETIC PROTEIN-16 (BMP-16)
TITLE OF INVENTION: COMPOSITIONS
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: GENETICS INSTITUTE, INC.
STREET: 87 CAMBRIDGE PARK DRIVE
CITY: CAMBRIDGE
STATE: MA
COUNTRY: USA
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/715 202A
FILING DATE: September 18, 1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: LAZAR, STEVEN R.
REGISTRATION NUMBER: 32,618
REFERENCE/DOCKET NUMBER: 5275
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8260
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 843 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: s19-peptide
LOCATION: 1..510
FEATURE:
NAME/KEY: mat-peptide
LOCATION: 511..840
FEATURE:
NAME/KEY: CDS
LOCATION: 1..840
US-08-715-202A-1
Query Match 83.2%; Score 15.8; DB 2; Length 843;
Best Local Similarity 89.5%; Pred. No. 20;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 ttgtctctccagaggtc 19
db 412 ttgtctctccagaggtt 394
RESULT 3
US-08-715-202A-3/c
Sequence 3, Application US/08715202A
Patent No. 5363403
GENERAL INFORMATION:
APPLICANT: CELESTE, ANTHONY J.
APPLICANT: MURRAY, BETH L.
TITLE OF INVENTION: BONE MORPHOGENETIC PROTEIN-16 (BMP-16)
TITLE OF INVENTION: COMPOSITIONS
NUMBER OF SEQUENCES: 10

CORRESPONDENCE ADDRESS:
ADDRESSEE: GENETICS INSTITUTE, INC.
STREET: 87 CAMBRIDGE PARK DRIVE
CITY: CAMBRIDGE
STATE: MA
COUNTRY: USA
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/715,202A
FILING DATE: September 18, 1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: LAZAR, STEVEN R.
REGISTRATION NUMBER: 32,618
REFERENCE/DOCKET NUMBER: 5275
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8260
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 2002 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-715-202A-3
Query Match 83.2%; Score 15.8; DB 2; Length 2002;
Best Local Similarity 89.5%; Pred. No. 23;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 ttgtctctccagaggtc 19
db 801 ttgtctctccagaggtt 783
RESULT 4
US-08-366-276-1
Sequence 1, Application US/08366276
Patent No. 5534409
GENERAL INFORMATION:
APPLICANT: Groner, Bernd
APPLICANT: Gouilleux, Fabrice
APPLICANT: Wakao, Hiroshi
TITLE OF INVENTION: cytokine Regulated Transcription Factor
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESSEE: CIBA-GEIGY Corporation
STREET: 7 Skyline drive
CITY: Hawthorne
STATE: New York
COUNTRY: USA
ZIP: 10532
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30B
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/366,276
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9409396.0
FILING DATE: 11-MAY-1994
ATTORNEY/AGENT INFORMATION:
NAME: Elmer, James Scott

REGISTRATION NUMBER: 36,129
REFERENCE/DOCKET NUMBER: 4-19992/A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (919) 541-8614
TELEFAX: (919) 541-8689
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2818 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 244..2625
OTHER INFORMATION: /product= "mammary gland factor"

Query Match 83.2%; Score 15.8; DB 1; Length 2818;
Best Local Similarity 89.5%; Pred. No. 24;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ttgtctctgcagaggtc 19
Db 2599 TTTACTCTGCGCAGAGCT 2617

RESULT 5
US-08-373-579-5
Sequence 5, Application US/08373579
Patent No. 5650490
GENERAL INFORMATION:
APPLICANT: DAVIS, et al.
TITLE OF INVENTION: TIE-2 LIGAND, METHOD OF MAKING AND USES
TITLE OF INVENTION: THEREOF
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Regeneron Pharmaceuticals, Inc.
STREET: 777 Old Saw Mill River Road
CITY: Tarrytown
STATE: New York
COUNTRY: USA
ZIP: 10591
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/373,579
FILING DATE: 17-JAN-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/353,503
FILING DATE: 09-DEC-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/348,492
FILING DATE: 02-DEC-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/330,261
FILING DATE: 27-OCT-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/319,932
FILING DATE: 07-OCT-1994
ATTORNEY/AGENT INFORMATION:
NAME: Cobert, Robert J.
REGISTRATION NUMBER: 36,108
REFERENCE/DOCKET NUMBER: REG 330-D
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 345-7400
TELEFAX: (914) 345-7721

INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 2282 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 357..1847
US-08-373-579-5

Query Match 77.9%; Score 14.8; DB 1; Length 2282;
Best Local Similarity 88.9%; Pred. No. 75;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ttgtctctgcagaggtc 18
Db 488 TTTCTCTGCGCAGAGAT 505

RESULT 6
US-08-418-595-5
Sequence 5, Application US/08418595
Patent No. 5814464
GENERAL INFORMATION:
APPLICANT: DAVIS, et al.
TITLE OF INVENTION: TIE-2 LIGAND, METHOD OF MAKING AND USES
TITLE OF INVENTION: THEREOF
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Regeneron Pharmaceuticals, Inc.
STREET: 777 Old Saw Mill River Road
CITY: Tarrytown
STATE: New York
COUNTRY: USA
ZIP: 10591
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/418,595
FILING DATE: 06-APR-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/373,579
FILING DATE: 17-JAN-1995
APPLICATION NUMBER: US 08/353,503
FILING DATE: 09-DEC-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/348,492
FILING DATE: 02-DEC-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/330,261
FILING DATE: 27-OCT-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/319,932
FILING DATE: 07-OCT-1994
ATTORNEY/AGENT INFORMATION:
NAME: Cobert, Robert J.
REGISTRATION NUMBER: 36,108
REFERENCE/DOCKET NUMBER: REG 330-D
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 345-7400
TELEFAX: (914) 345-7721
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 2282 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 357..1847
US-08-418-595-5

Query Match 77.9%; Score 14.8; DB 1; Length 2282;
Best Local Similarity 88.9%; Pred. No. 75;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ttgtctctgcagaggt 18
||| |||||
DB 488 ttctctctgcagagat 505

RESULT 7
US-08-665-926-5
Sequence 5, Application US/08665926
Patent No. 5851797
GENERAL INFORMATION:
APPLICANT: Valenzuela et al.
TITLE OF INVENTION: THE LIGAND-3, METHODS OF MAKING AND USES
TITLE OF INVENTION: THEREOF
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: Regeneron Pharmaceuticals, Inc.
STREET: 777 Old Saw Mill River Road
CITY: Tarrytown
STATE: New York
COUNTRY: U.S.A.
ZIP: 10591-6707
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/665,926
FILING DATE: 19-JUN-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Robert J. Cobert
REGISTRATION NUMBER: 36,108
REFERENCE/DOCKET NUMBER: REG 330-H
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 345-7400
TELEFAX: (914) 345-2113
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 2282 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 357..1847
US-08-665-926-5

Query Match 77.9%; Score 14.8; DB 2; Length 2282;
Best Local Similarity 88.9%; Pred. No. 75;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 ttgtctctgcagaggt 18
||| |||||
DB 488 ttctctctgcagagat 505

RESULT 8
US-09-162-437-5

Sequence 5, Application US/09162437
Patent No. 6166185

GENERAL INFORMATION:
APPLICANT: Davis, et al.
TITLE OF INVENTION: TIF-2 LIGAND, METHOD OF MAKING AND USES
TITLE OF INVENTION: THEREOF
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Regeneron Pharmaceuticals, Inc.
STREET: 777 Old Saw Mill River Road
CITY: Tarrytown
STATE: New York
COUNTRY: USA
ZIP: 10591

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/162,437
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/418,595
FILING DATE: 06-APR-1995

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/373,579
FILING DATE: 17-JAN-1995

APPLICATION NUMBER: US 08/353,503
FILING DATE: 09-DEC-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/348,492
FILING DATE: 02-DEC-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/330,261
FILING DATE: 27-OCT-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/319,932
FILING DATE: 07-OCT-1994

ATTORNEY/AGENT INFORMATION:
NAME: Cobert, Robert J.
REGISTRATION NUMBER: 36,108
REFERENCE/DOCKET NUMBER: REG 330-D
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 345-7400
TELEFAX: (914) 345-7721
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 2282 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 357..1847
US-09-162-437-5

Query Match 77.9%; Score 14.8; DB 4; Length 2282;
Best Local Similarity 88.9%; Pred. No. 75;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 ttgtctctgcagaggt 18
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DB 488 ttctctctgcagagat 505

RESULT 9
US-08-447-411-44
Sequence 44, Application US/08447411
Patent No. 5773243

GENERAL INFORMATION:
APPLICANT: FRITZINGER, DAVID C.
APPLICANT: BREDEHORST, REINHARD
APPLICANT: VOGEL, CARL-WILHELM
TITLE OF INVENTION: DNA ENCODING COBRA C3, CVF1, AND CVF2
NUMBER OF SEQUENCES: 81
CORRESPONDENCE ADDRESS:
ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
ADDRESS: P.C.
STREET: 1755 S. Jefferson Davis Highway, Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/447,411
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/043,747
FILING DATE: 07-APR-1993
ATTORNEY/AGENT INFORMATION:
NAME: Oblon, NO. 5773243man F.
REGISTRATION NUMBER: 24,618
REFERENCE/DOCKET NUMBER: 1126-101-0
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 413-3000
TELEFAX: (703) 413-2220
TELEX: 248355 OPAT UR
INFORMATION FOR SEQ. ID NO: 44:
SEQUENCE CHARACTERISTICS:
LENGTH: 5924 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: sig-peptide
LOCATION: 4..69
FEATURE:
NAME/KEY: mat-peptide
LOCATION: 70..4929
FEATURE:
NAME/KEY: CDS
LOCATION: 4..4929
US-08-447-411-44

Query Match 77.9%; Score 14.8; DB 1; Length 5924;
Best Local Similarity 88.9%; Pred. No. 87;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ttgtctctgcacagagt 18
|| ||||| |||||
Db 5567 TTGGCTCTCTGCACAGAGT 5564

RESULT 10
US-08-662-227-1
Sequence 1, Application US/08662227
Patent No. 5923320
GENERAL INFORMATION:
APPLICANT: VOGEL, CARL-WILHELM
APPLICANT: BREDEHORST, REINHARD
APPLICANT: KOCK, MICHAEL
APPLICANT: FRITZINGER, DAVID
TITLE OF INVENTION: RECOMBINANT PROCVF
NUMBER OF SEQUENCES: 39

CORRESPONDENCE ADDRESS:
ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
ADDRESS: P.C.
STREET: 1755 S. JEFFERSON DAVIS HIGHWAY
CITY: ARLINGTON
STATE: VA
COUNTRY: USA
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/662,227
FILING DATE: 14-JUN-1996
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: OBLON, NORMAN F.
REGISTRATION NUMBER: 24,618
REFERENCE/DOCKET NUMBER: 1126-0107-0X
TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-413-3000
TELEFAX: 703-413-2220
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5948 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-662-227-1

Query Match 77.9%; Score 14.8; DB 2; Length 5948;
Best Local Similarity 88.9%; Pred. No. 87;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ttgtctctgcacagagt 18
|| ||||| |||||
Db 5567 TTGGCTCTCTGCACAGAGT 5564

RESULT 11
US-08-359-705B-7/C
Sequence 7, Application US/08359705B
Patent No. 5844092
GENERAL INFORMATION:
APPLICANT: Presta, Leonard G.
APPLICANT: Shelton, David L.
APPLICANT: Ufer, Roman
TITLE OF INVENTION: Human trk Receptors and Neurotrophic Factor Inhibitors
NUMBER OF SEQUENCES: 41
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genentech, Inc.
STREET: 1 DNA Way
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: WinpatIn (Genentech)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/359,705B
FILING DATE: 20-Dec-1994
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/286646
FILING DATE: 08/10/94
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/215139

FILING DATE: 03/18/94

ATTORNEY/AGENT INFORMATION:

NAME: TORCHLA, PHD., TIMOTHY E.

REGISTRATION NUMBER: 36,700

REFERENCE/DOCKET NUMBER: P0873P2

TELECOMMUNICATION INFORMATION:

TELEPHONE: 650/225-8674

TELEFAX: 650/952-9861

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1858 base pairs

TYPE: Nucleic Acid

STRANDEDNESS: Single

TOPOLOGY: Linear

US-08-359-7058-7

Query Match

Best Local Similarity 75.8%; Score 14.4; DB 2; Length 1858;

Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 tgcctctgccagaggt 18

DB 545 TGCTCTGCCAGAGCT 530

RESULT 12

US-08-286-846A-7/C

Sequence 7, Application US/08286846A

Patent No. 5877016

GENERAL INFORMATION:

APPLICANT: Prestia, Leonard G.

APPLICANT: Shelton, David L.

TITLE OF INVENTION: Human trk Receptors and Neurotrophic Factor Inhibitors

NUMBER OF SEQUENCES: 41

CORRESPONDENCE ADDRESS:

ADDRESSER: Genentech, Inc.

STREET: 460 Point San Bruno Blvd

CITY: South San Francisco

STATE: California

COUNTRY: USA

ZIP: 94080

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk

OPERATING SYSTEM: IBM PC compatible

SOFTWARE: Winpatin (Genentech)

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/286,846A

FILING DATE: 05-Aug-1994

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: TORCHLA, PHD., TIMOTHY E.

REGISTRATION NUMBER: 36,700

REFERENCE/DOCKET NUMBER: P0873P1

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415/225-8674

TELEFAX: 415/952-9861

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1858 base pairs

TYPE: Nucleic Acid

STRANDEDNESS: Single

TOPOLOGY: Linear

US-08-286-846A-7

QY 3 tgcctctgccagaggt 18

DB 545 TGCTCTGCCAGAGCT 530

RESULT 13

US-08-457-880A-7/C

Sequence 7, Application US/08457880A

Patent No. 5910574

GENERAL INFORMATION:

APPLICANT: Leonard G. Prestia

APPLICANT: David L. Shelton

TITLE OF INVENTION: HUMAN trk RECEPTORS AND NEUROTROPHIC FACTOR

NUMBER OF SEQUENCES: 41

CORRESPONDENCE ADDRESS:

ADDRESSER: Genentech, Inc.

STREET: 1 DNA Way

CITY: South San Francisco

STATE: California

COUNTRY: USA

ZIP: 94080

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk

OPERATING SYSTEM: IBM PC compatible

SOFTWARE: Winpatin (Genentech)

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/457,880A

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/444,622

FILING DATE: 19-May-1995

APPLICATION NUMBER: 08/286846

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: TORCHLA, PHD., TIMOTHY E.

REGISTRATION NUMBER: 36,700

REFERENCE/DOCKET NUMBER: P0873P1C3

TELECOMMUNICATION INFORMATION:

TELEPHONE: 650/225-8674

TELEFAX: 650/952-9861

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1858 base pairs

TYPE: Nucleic Acid

STRANDEDNESS: Single

TOPOLOGY: Linear

US-08-457-880A-7

Query Match

Best Local Similarity 75.8%; Score 14.4; DB 2; Length 1858;

Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 tgcctctgccagaggt 18

DB 545 TGCTCTGCCAGAGCT 530

RESULT 14

US-08-444-622A-7/C

Sequence 7, Application US/08444622A

Patent No. 6025166

GENERAL INFORMATION:

APPLICANT: Leonard G. Prestia

APPLICANT: David L. Shelton

TITLE OF INVENTION: HUMAN trk RECEPTORS AND NEUROTROPHIC FACTOR INHIBITORS

NUMBER OF SEQUENCES: 41

CORRESPONDENCE ADDRESS:

ADDRESSEE: Genentech, Inc.
STREET: 1 DNA Way
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Minipalin (Genentech)

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/444,622A
FILING DATE: 19-May-1995

CLASSIFICATION: 424

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/286846

FILING DATE: 5

ATTORNEY/AGENT INFORMATION:

NAME: Torchia, PhD., Timothy E.
REGISTRATION NUMBER: 36,700

REFERENCE/DOCKET NUMBER: P0873PIC3
TELECOMMUNICATION INFORMATION:

TELEPHONE: 650/225-8674
TELEFAX: 650/952-9881

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1858 base pairs
TYPE: Nucleic Acid

STRANDEDNESS: Single
TOPOLOGY: Linear

US-08-444-622A-7

Query Match

Best Local Similarity 75.8%; Score 14.4; DB 3; Length 1858;

Matches 15; Conservativity 93.8%; Pred. No. 1.2e+02;

Mismatches 0; Indels 1; Gaps 0;

QY 3 tgcctctgccagaggt 18
|||||

DB 545 TGCCTCTGCCAGAGCT 530

RESULT 15

US-08-942-562-7/C

Sequence 7, Application US/08942562

Patent No. 6027927

GENERAL INFORMATION:

APPLICANT: Presta, Leonard G.
APPLICANT: Shelton, David L.

APPLICANT: Uffler, Roman

TITLE OF INVENTION: Human trk Receptors and Neurotrophic

NUMBER OF SEQUENCES: 37

CORRESPONDENCE ADDRESS:

ADDRESSEE: Genentech, Inc.
STREET: 460 Point San Bruno Blvd

CITY: South San Francisco
STATE: California

COUNTRY: USA
ZIP: 94080

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Minipalin (Genentech)

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/942,562

FILING DATE: 01-OCT-1997

CLASSIFICATION: 530

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/444,597

FILING DATE: 19-May-1995

ATTORNEY/AGENT INFORMATION:

NAME: Torchia, PhD., Timothy E.
REGISTRATION NUMBER: 36,700

REFERENCE/DOCKET NUMBER: P0873PIC2
TELECOMMUNICATION INFORMATION:

TELEPHONE: 415/225-8674
TELEFAX: 415/952-9881

TELEX: 910/371-7168

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1858 base pairs
TYPE: Nucleic Acid

STRANDEDNESS: Single
TOPOLOGY: Linear

US-08-942-562-7

Query Match

Best Local Similarity 75.8%; Score 14.4; DB 3; Length 1858;

Matches 15; Conservativity 93.8%; Pred. No. 1.2e+02;

Mismatches 0; Indels 1; Gaps 0;

QY 3 tgcctctgccagaggt 18
|||||

DB 545 TGCCTCTGCCAGAGCT 530

Search completed: July 25, 2001, 05:18:55
Job time: 9218 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sv model

Run on: July 25, 2001, 02:01:07 ; Search time 2762.24 Seconds

(without alignments)
71.866 Million cell updates/sec

Title: US-09-142-095-1

Perfect score: 21

Sequence: 1 aagtgacctccctgctacctt 21

Scoring table: IDENTITY_NDC
Gapop 10.0 , Gapext 1.0

Searched: 10228115 seqs, 4726426750 residues

Total number of hits satisfying chosen parameters: 20456230

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	17.8	84.8	212	R68057	R68057 yH98612.r1
2	17.8	84.8	348	R31201	R31201 yH62802.r1
3	17.8	84.8	417	R67982	R67982 y10410.r1
4	17.8	84.8	418	R31714	R31714 yH6312.r1
5	17.8	84.8	481	R63154	R63154 y10111.r1
6	17.8	84.8	498	225	A0882343 HS-5382.A
7	17.8	84.8	510	225	A0185331 HS-3065.A
8	17.8	84.8	608	230	A051067 RPCI-11-4
9	17.8	81.0	471	31	AV553454 AV553454
10	16.8	80.0	193	115	AM381670 QVO-57031
11	16.8	80.0	304	8	AA500281 V197108.r
12	16.8	80.0	326	9	AA560193 V120501.r
13	16.8	80.0	410	9	AA620205 V064806.r
14	16.8	80.0	416	22	A1615361 V197108.y
15	16.8	80.0	439	9	AA616429 V069602.r
16	16.8	80.0	469	8	AA473687 V899605.x
17	16.8	80.0	508	13	AA881891 Vx2910.r
18	16.8	80.0	514	236	A0978201 RPCI-23-2
19	16.8	80.0	535	238	A2093040 RPCI-23-4
20	16.8	80.0	547	120	AAW43388 up84h01.y
21	16.8	80.0	556	2	AA108719 mp29f06.r
22	16.8	80.0	561	238	A2123144 RPCI-23-4
23	16.8	80.0	740	245	A2616170 RPCI-23-1
24	16.4	78.1	322	251	A2902873 RPCI-24-1
25	16.4	78.1	415	147	BF370208 RCI-FR003
26	16.4	78.1	481	114	AW342357 G1HE57414
27	16.4	78.1	508	21	A149582 tn96e01.x
28	16.4	78.1	530	123	AM981762 PC18D09.P
29	16.4	78.1	665	240	A263471 RPCI-23-1
30	16.4	78.1	668	246	A2616170 RCI-23-1
31	16.4	78.1	783	155	BC573720 602594785
32	16.4	78.1	810	221	CNS03GRY
33	16.4	78.1	911	147	BF304665
34	16.4	78.1	914	220	CNS01RCS
35	16.4	78.1	1049	222	CNS05R8X
36	16.2	77.1	193	218	AF149685
37	16.2	77.1	265	11	AA716271 zH28611.s
38	16.2	77.1	274	23	A1651623 wD07d07.x
39	16.2	77.1	285	15	A1021546 uD09c11.r
40	16.2	77.1	319	14	AA985674 or71e07.s
41	16.2	77.1	340	24	A1744438 wF89c03.x
42	16.2	77.1	351	4	AA235220 z836e12.r
43	16.2	77.1	355	123	AM999271 MR0-BN007
44	16.2	77.1	370	18	A1274461
45	16.2	77.1	378	168	BF722600 mab21c09.

ALIGNMENTS

RESULT	1	212 bp	EST	01-JUN-1995
LOCUS	R68057	212 bp	EST	01-JUN-1995
DEFINITION	yH98612.r1 Soares placenta NB2HP Homo sapiens cDNA clone			
IMAGE	137806 5', mRNA sequence.			
ACCESSION	R68057.1	GI:841574		
VERSION	R68057.1	GI:841574		
KEYWORDS	EST.			
SOURCE	human.			
ORGANISM	Homo sapiens			
REFERENCE	1 (bases 1 to 212)			
AUTHORS	Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Maitre, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston, R., Williamson, A., Wohldmann, P. and Wilson, R.			

TITLE

The WashU-Merck EST Project
Unpublished (1995)

JOURNAL

Contact: Wilson RK

COMMENT

Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu

FEATURES

High quality sequence stops: 74
Insert size: 761
Source: IMAGE Consortium, LNL
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert Length: 761 Std Error: 0.00
Seq primer: M13P1
High quality sequence stop: 74.

FEATURES

Location/Qualifiers
1..212
/organism="Homo sapiens"
/db_xref="GDB:544134"
/db_xref="taxon:9606"
/clone="IMAGE:137806"
/clone.lib="Soares placenta NB2HP"
/sex="Female"
/dev_stage="placenta obtained at birth (full term)"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: placenta; Vector: pTZ19 (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer (5'
AAGCGAAGATTCGGCGCCGACGAGATTTTCTTTTCTTTT 3')
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pTZ19 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M. Patricia Bonafino."

BASE COUNT

50 a 63 c 32 g 66 t 1 others

Query Match

84.8%; Score 17.8; DB 188; Length 212;
Best Local Similarity 90.5%; Pred. No. 1.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

DB

11 AAGTACCTCCGCTACCTT 31

RESULT

2
R31201 348 bp mRNA EST 28-APR-1995

LOCUS

yH62802.r1 Soares placenta NB2HP Homo sapiens cDNA clone

DEFINITION

IMAGE:134282 5' similar to contains MSRI repetitive element ; mRNA
sequence.

ACCESSION

R31201.1 GI:787044

VERSION

R31201

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

REFERENCE

1 (bases 1 to 348)

AUTHORS

Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Maitre, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston, R., Williamson, A., Wohldmann, P. and Wilson, R.

TITLE

Unpublished (1995)

JOURNAL

Contact: Wilson RK

COMMENT

Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Email: estevason.wustl.edu

Insert Size: 728

High quality sequence stops: 195

Source: IMAGE Consortium, LNL

This clone is available royalty-free through LNL; contact the

IMAGE Consortium (info@image.lnl.gov) for further information.

Insert Length: 728

Seq primer: M13RPI

Std Error: 0.00

High quality sequence stop: 195.

Location/Qualifiers

FEATURES

source

1. 348

/organism="Homo sapiens"

/db_xref="GDB:540030"

/db_xref="taxon:9606"

/clone="IMAGE:134282"

/clone_lib="Soares placenta Nb2HP"

/sex="Female"

/dev_stage="Placenta obtained at birth (full term)"

/lab_host="DH10B (ampicillin resistant)"

/note="Organ: placenta; Vector: pT73D (Pharmacia) with a

modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st

strand cDNA was primed with a Not I - oligo(dT) primer [5'

AACTGAGAGATTCGGCGCGGAGATTTTCTTTTCTTTT 3'],

double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Not I and cloned into the Not I

and Eco RI sites of the modified pT73 vector. Library

went through one round of normalization. Library

constructed by Bento Soares and M. Fatima Bonaldo.

BASE COUNT

78 a 92 c 66 g 108 t 4 others

ORIGIN

Query Match 84.8%; Score 17.8; DB 187; Length 348;

Best Local Similarity 90.5%; Pred. No. 1.5e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 aagtgactccctgactct 21

||||| |||||||

Db 10 AAGTAGCTCCCTGCTACTT 30

RESULT 3

LOCUS

R67982

DEFINITION

Y104410.r1 Soares placenta Nb2HP Homo sapiens cDNA clone

IMAGE:138234 5', mRNA sequence.

ACCESSION

R67982

VERSION

R67982.1

KEYWORDS

EST.

SOURCE

HUMAN.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 417)

AUTHORS

Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman

, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Maita, M., Parsons, J.,

Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston

, R., Williamson, A., Wohlmann, P. and Wilson, R.

The Mashu-Merck EST Project

Unpublished (1995)

TITLE

JOURNAL

COMMENT

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: estevason.wustl.edu

Insert Size: 774

High quality sequence stops: 306

Source: IMAGE Consortium, LNL

This clone is available royalty-free through LNL; contact the

IMAGE Consortium (info@image.lnl.gov) for further information.

Insert Length: 774

Std Error: 0.00

Seq primer: M13RPI

FEATURES

source

1. 417

High quality sequence stop: 306.

Location/Qualifiers

/organism="Homo sapiens"

/db_xref="GDB:544609"

/db_xref="taxon:9606"

/clone="IMAGE:138234"

/clone_lib="Soares placenta Nb2HP"

/sex="Female"

/dev_stage="Placenta obtained at birth (full term)"

/lab_host="DH10B (ampicillin resistant)"

/note="Organ: placenta; Vector: pT73D (Pharmacia) with a

modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st

strand cDNA was primed with a Not I - oligo(dT) primer [5'

AACTGAGAGATTCGGCGCGGAGATTTTCTTTTCTTTT 3'],

double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Not I and cloned into the Not I

and Eco RI sites of the modified pT73 vector. Library

went through one round of normalization. Library

constructed by Bento Soares and M. Fatima Bonaldo.

BASE COUNT

104 a 101 c 89 g 123 t

ORIGIN

Query Match 84.8%; Score 17.8; DB 188; Length 417;

Best Local Similarity 90.5%; Pred. No. 1.6e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 aagtgactccctgactct 21

||||| |||||||

Db 10 AAGTAGCTCCCTGCTACTT 30

RESULT 4

LOCUS

R31714

DEFINITION

Yh63a12.r1 Soares placenta Nb2HP Homo sapiens cDNA clone

IMAGE:134398 5', mRNA sequence.

ACCESSION

R31714

VERSION

R31714.1

KEYWORDS

EST.

SOURCE

HUMAN.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 418)

AUTHORS

Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman

, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Maita, M., Parsons, J.,

Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston

, R., Williamson, A., Wohlmann, P. and Wilson, R.

The Mashu-Merck EST Project

Unpublished (1995)

TITLE

JOURNAL

COMMENT

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: estevason.wustl.edu

Insert Size: 707

High quality sequence stops: 244

Source: IMAGE Consortium, LNL

This clone is available royalty-free through LNL; contact the

IMAGE Consortium (info@image.lnl.gov) for further information.

Insert Length: 707

Std Error: 0.00

Seq primer: M13RPI

High quality sequence stop: 244.

Location/Qualifiers

1. 418

/organism="Homo sapiens"

/db_xref="GDB:540219"

/db_xref="taxon:9606"

/clone="IMAGE:134398"

/clone_lib="Soares placenta Nb2HP"

/sex="Female"
 /dev-stage="placenta obtained at birth (full term)"
 /lab_host="DH10B (ampicillin resistant)"
 /note="Organ: Placenta; Vector: pT73D (Pharmacia) with a modified polylinker; Site: 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' AACGAGAGAAATTCGCGCCGACAGAAATTTTCTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."
 BASE COUNT 90 a 100 c 95 g 126 t 7 others
 ORIGIN

Query Match 84.8%; Score 17.8; DB 187; Length 418;
 Best Local Similarity 90.5%; Pred. No. 1.6e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 aagtgactccctgctaccc 21
 Db 10 AAGTTAGCTCCCTGCTACCTT 30

RESULT 5
 R63154 481 bp mRNA EST 26-MAY-1995
 LOCUS Y101a11.r1 Soares Placenta Nb2HP Homo sapiens cDNA clone
 DEFINITION IMAGE:137948 5', mRNA sequence.
 ACCESSION R63154
 VERSION R63154.1 GI:835033
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 481)
 AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Merra, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.
 COMMENT The Washu-Merck EST Project
 JOURNAL Unpublished (1995)
 MEDLINE
 COMMENT Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@wustl.wustl.edu
 Insert Size: 769
 High quality sequence stops: 411
 Source: IMAGE Consortium, LNL
 This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
 Seg primer: M13RPI
 High quality sequence stop: 411.
 Location/Qualifiers

FEATURES

source

1. 481
 /organism="Homo sapiens"
 /db_xref="GDB:544291"
 /db_xref="taxon:9606"
 /clone="IMAGE:137948"
 /clone_lib="Soares Placenta Nb2HP"
 /sex="Female"
 /dev-stage="placenta obtained at birth (full term)"
 /lab_host="DH10B (ampicillin resistant)"
 /note="Organ: Placenta; Vector: pT73D (Pharmacia) with a modified polylinker; Site: 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' AACGAGAGAAATTCGCGCCGACAGAAATTTTCTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."
 BASE COUNT 112 a 114 c 99 g 152 t 4 others
 ORIGIN

Query Match 84.8%; Score 17.8; DB 188; Length 481;
 Best Local Similarity 90.5%; Pred. No. 1.6e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 aagtgactccctgctaccc 21
 Db 11 AAGTTAGCTCCCTGCTACCTT 31

RESULT 6
 A0882343 498 bp DNA GSS 09-NOV-1999
 LOCUS HS_5382.A1-B03-T7C RPCI-11 Human Male BAC Library Homo sapiens
 DEFINITION genomic clone Plate-9150 Col-5 Row-C, DNA sequence.
 ACCESSION A0882343
 VERSION A0882343.1 GI:6313810
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 498)
 AUTHORS Mahairas, G.G., Wallace, J.C., Smith, R., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and Hood, L.
 COMMENT Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
 Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
 JOURNAL 99380589
 MEDLINE
 COMMENT Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@u.washington.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.bufileo.edu/ordering/bac.htm>) or from Research Genetics (<http://www.resgen.com>). BAC end Web Server: <http://www.hsc.washington.edu>
 Plate: 9150 row: C column: 5
 Seg primer: T7
 Class: BAC ends
 High quality sequence stop: 498.
 Location/Qualifiers

FEATURES

source

1. 498
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Plate-9150 Col-5 Row-C"
 /clone_lib="RPCI-11 Human Male BAC Library"
 /sex="male"
 /note="Vector: pBAC3.6; Site: 1: EcoRI; Site 2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"
 BASE COUNT 155 a 111 c 118 g 111 t 3 others
 ORIGIN

Query Match 84.8%; Score 17.8; DB 235; Length 498;
 Best Local Similarity 90.5%; Pred. No. 1.6e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

ORGANISM	Homo sapiens
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE	1 (Pages 1 to 608)
AUTHORS	Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter

REFERENCE	AUTHORS	TITLE	JOURNAL	MEDLINE	COMMENT
1 (pages 1 to 471)	Asamizu, E., Nakamura, Y., Sato, S. and Tabata, S.	A large scale analysis of cDNA in Arabidopsis thaliana: Generation of 12,028 non-redundant expressed sequence tags from normalized and size-selected cDNA libraries	DNA Res. 7, 175-180 (2000)	20363093	Contact: Erika Asamizu The First Laboratory for Plant Gene Research Kazusa DNA Research Institute Yama 1532-3, Kisarazu, Chiba 292-0812, Japan Email: asamizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/location/Qualifiers

FEATURES	Location/Qualifiers
source	I..471
	/organism="Arabidopsis thaliana"
	/strain="Columbia"
	/db_xref="taxon:3702"

RESULT	12
AA560193	
LOCUS	
DEFINITION	AA560193 326 bp mRNA EST 18-AUG-1997
ACCESSION	V120901.r1 StrataGene mouse Tcell1 937311 Mus musculus cDNA clone
VERSION	AA560193 IMAGE:772616 5', mRNA sequence.
KEYWORDS	AA560193.1 GI 2331656
	EST.

COMMENT
Contact: Maria Wasmu-NCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8301, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1800
Email: mouseest@wustl.edu
This clone is available royalty-free through LNM; contact the

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using SW model

Run on: July 25, 2001, 05:18:51 ; Search time 117.39 Seconds
(without alignments)
33.141 Million cell updates/sec

Title: US-09-142-095-2

Perfect score: 21
Sequence: 1 ccactgggatacaagatctc 21

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 317530 seqs, 92630169 residues

Total number of hits satisfying chosen parameters: 635060

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

Database :

Issued Patents.NA.*
1: /cgn2_6/pdata/2/1na/5a_COMB.seq:*
2: /cgn2_6/pdata/2/1na/5b_COMB.seq:*
3: /cgn2_6/pdata/2/1na/6a_COMB.seq:*
4: /cgn2_6/pdata/2/1na/6b_COMB.seq:*
5: /cgn2_6/pdata/2/1na/PCPUS_COMB.seq:*
6: /cgn2_6/pdata/2/1na/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match Length	DB ID	Description
C 1	21	100.0	1190	5	PCT-US92-00282-18
C 2	21	100.0	2336	5	PCT-US97-00282-1
C 3	15.8	75.2	1073	4	US-08-960-780-37
C 4	15.8	75.2	1073	4	US-08-073-898-37
C 5	14.8	70.5	831	2	US-08-403-853-17
C 6	14.8	70.5	832	1	US-08-473-496-1
C 7	14.8	70.5	832	2	US-08-454-028-3
C 8	14.8	70.5	832	5	PCT-US94-05388-3
C 9	14.8	70.5	832	5	PCT-US95-09121-1
C 10	14.8	70.5	832	5	PCT-US96-07496-3
C 11	14.8	70.5	3225	1	US-08-698-551-13
C 12	14.8	70.5	3225	2	US-08-602-228-13
C 13	14.8	70.5	3225	2	US-08-533-901B-13
C 14	14.8	70.5	3225	2	US-08-839-032A-13
C 15	14.8	70.5	3225	2	US-08-839-032A-13
C 16	14.8	70.5	3225	5	PCT-US95-12724-13
C 17	14.6	69.5	87350	3	US-08-781-891-79
C 18	14.2	67.6	20	4	US-08-669-378-16
C 19	14.2	67.6	330	1	US-08-558-735-19
C 20	14.2	67.6	330	4	US-08-906-480-19
C 21	14.2	67.6	333	1	US-08-558-735-18
C 22	14.2	67.6	333	4	US-08-906-480-17
C 23	14.2	67.6	336	2	US-08-888-366-17
C 24	14.2	67.6	343	3	US-08-320-559-36
C 25	14.2	67.6	343	3	US-08-545-860D-36
C 26	14.2	67.6	343	5	PCT-US94-04496-36
C 27	14.2	67.6	345	1	US-08-558-735-3

C 28	14.2	67.6	345	1	US-08-558-735-17	Sequence 17, Appl
C 29	14.2	67.6	345	4	US-08-906-480-3	Sequence 3, Appl
C 30	14.2	67.6	345	4	US-08-906-480-17	Sequence 17, Appl
C 31	14.2	67.6	355	1	US-08-558-735-16	Sequence 16, Appl
C 32	14.2	67.6	355	4	US-08-906-480-16	Sequence 16, Appl
C 33	14.2	67.6	428	3	US-08-589-939-4	Sequence 4, Appl
C 34	14.2	67.6	532	1	US-08-558-735-1	Sequence 1, Appl
C 35	14.2	67.6	532	4	US-08-906-480-1	Sequence 1, Appl
C 36	14.2	67.6	780	3	US-09-027-449-54	Sequence 54, Appl
C 37	14.2	67.6	780	3	US-09-027-449-58	Sequence 54, Appl
C 38	14.2	67.6	780	4	US-09-027-449-65	Sequence 54, Appl
C 39	14.2	67.6	780	4	US-08-804-444A-54	Sequence 54, Appl
C 40	14.2	67.6	780	4	US-08-804-444A-58	Sequence 58, Appl
C 41	14.2	67.6	780	4	US-09-026-985-54	Sequence 54, Appl
C 42	14.2	67.6	780	4	US-09-026-985-58	Sequence 54, Appl
C 43	14.2	67.6	780	4	US-09-026-985-65	Sequence 54, Appl
C 44	14.2	67.6	1422	4	US-08-387-117-8	Sequence 8, Appl
C 45	14.2	67.6	1905	4	US-08-387-117-7	Sequence 7, Appl

ALIGNMENTS

RESULT 1
PCT-US92-00282-18/C
Sequence 18, Application PC/TUS9200282
GENERAL INFORMATION:
APPLICANT: OMENS, IDA S.
APPLICANT: RITTER, JOSEPH K.
TITLE OF INVENTION: THE GENETIC LOCUS UGT1 AND A MUTATION
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: CUSHMAN DABRY & CUSHMAN
STREET: 1615 L STREET, N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20036-5601
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/00282
FILING DATE: 19920110
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: SCOTT, WATSON T.
REGISTRATION NUMBER: 26581
REFERENCE/DOCKET NUMBER: 91532-PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-861-3000
TELEFAX: 202-822-0944
TELEX: 6714627 CUSH
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 1190 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
PCT-US92-00282-18
Query Match 100.0%; Score 21; DB 5; Length 1190;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1 ccactgggatacaagatctc 21
DB 190 CCACCTGGATCAACAGATCTC 170

RESULT 2
PCT-US92-00282-1/c
Sequence 1, Application PC/TUS9200282
GENERAL INFORMATION:
APPLICANT: OMENS, IDA S.
TITLE OF INVENTION: THE GENETIC LOCUS UGT1 AND A MUTATION
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: CUSHMAN DABBY & CUSHMAN
STREET: 1615 L STREET, N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20036-5601
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/00282
FILING DATE: 19920110
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: SCOTT, WATSON T.
REGISTRATION NUMBER: 26581
REFERENCE/DOCKET NUMBER: 91532-PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-861-3000
TELEFAX: 202-822-0944
TELEX: 6714627 CUSH
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2336 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
PCT-US92-00282-1

Query Match 100.0%; Score 21; DB 5; Length 2336;
Best Local Similarity 100.0%; Pred. No. 0.082;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ccaactggatcacacagatct 21
DB 106 CCACGTGGATCAACAGTACT 86

RESULT 3
US-08-960-780-37
Sequence 37, Application US/08960780
Patent No. 6204435
GENERAL INFORMATION:
APPLICANT: Feltelson, Jerald S.
APPLICANT: Schnepf, H. Ernest
APPLICANT: Narva, Kenneth E.
APPLICANT: Stockhoff, Brian A.
APPLICANT: Schmeltz, James
APPLICANT: Loewer, David
APPLICANT: Dullum, Charles Joseph
APPLICANT: Muller-Cohn, Judy
APPLICANT: Stamp, Lisa
TITLE OF INVENTION: No. 6204435el Pesticidal Toxins and Nucleotide
TITLE OF INVENTION: Sequences Which Encode These Toxins
NUMBER OF SEQUENCES: 134
CORRESPONDENCE ADDRESS:
ADDRESSEE: Saliwanchik, Lloyd & Saliwanchik

STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: FL
COUNTRY: US
ZIP: 32606-6669
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/960,780
FILING DATE: 30-OCT-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/029,848
FILING DATE: 30-OCT-1996
ATTORNEY/AGENT INFORMATION:
NAME: Saliwanchik, David R.
REGISTRATION NUMBER: 31,794
REFERENCE/DOCKET NUMBER: MA-708
TELECOMMUNICATION INFORMATION:
TELEPHONE: 352-375-8100
TELEFAX: 352-372-5800
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 1073 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: 196F3
US-08-960-780-37

Query Match 75.2%; Score 15.8; DB 4; Length 1073;
Best Local Similarity 89.5%; Pred. No. .29;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcacacagatct 21
DB 838 ACTGGCAAAAACAGTACT 856

RESULT 4
US-09-073-898-37
Sequence 37, Application US/09073898
Patent No. 6242669
GENERAL INFORMATION:
APPLICANT: Feltelson, Jerald S.
APPLICANT: Schnepf, H. Ernest
APPLICANT: Narva, Kenneth E.
APPLICANT: Stockhoff, Brian A.
APPLICANT: Schmeltz, James
APPLICANT: Loewer, David
APPLICANT: Dullum, Charles Joseph
APPLICANT: Muller-Cohn, Judy
APPLICANT: Stamp, Lisa
APPLICANT: Morrill, George
APPLICANT: Flinstad-Lee, Stacey
TITLE OF INVENTION: No. 6242669el Pesticidal Toxins and Nucleotide
TITLE OF INVENTION: Sequences Which Encode These Toxins
NUMBER OF SEQUENCES: 144
CORRESPONDENCE ADDRESS:
ADDRESSEE: Saliwanchik, Lloyd & Saliwanchik
STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: FL
COUNTRY: US
ZIP: 32606-6669
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/073,898
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/029,848
FILING DATE: 30-OCT-1996
PRIOR APPLICATION DATA: US 08/960,780
APPLICATION NUMBER: US 08/960,780
FILING DATE: 30-OCT-1997
ATTORNEY/AGENT INFORMATION:
NAME: Sanders, Jay M.
REGISTRATION NUMBER: 39,355
REFERENCE/DOCKET NUMBER: MA-708C1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 352-375-8100
TELEFAX: 352-372-5800
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 1073 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: 196F3
US-09-073-898-37

Query Match 75.2% Score 15.8; DB 4; Length 1073;
Best Local Similarity 89.5%; Pred. No. 29;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcaacagatctc 21
||||| |||||||
DB 838 ACTGGGMAAACAGTATCT 856

RESULT 5
US-08-403-853-17
Sequence 17, Application US/08403853
Patent No. 5844094
GENERAL INFORMATION:
APPLICANT: HUDSON, Peter J.
APPLICANT: LAH, Maria
APPLICANT: KORST, Alex A.
APPLICANT: IRVING, Robert A.
APPLICANT: ATWELL, John L.
APPLICANT: MALBY, Robyn L.
APPLICANT: POWER, Barbara E.
APPLICANT: COLMAN, Peter M.
TITLE OF INVENTION: TARGET BINDING POLYPEPTIDE
NUMBER OF SEQUENCES: 25
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/403,853
FILING DATE: 30-MAY-1995
CLASSIFICATION: 433
PRIOR APPLICATION DATA:

APPLICATION NUMBER: WO PCT/AU93/00491
FILING DATE: 24-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: AU PL 4973
FILING DATE: 25-SEP-1992
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16786/189/CHAC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 831 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 1..819
FEATURE:
NAME/KEY: CDS
LOCATION: 1..819
US-08-403-853-17

Query Match 70.5% Score 14.8; DB 2; Length 831;
Best Local Similarity 88.9%; Pred. No. 88;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcaacagatc 20
||||| |||||||
DB 575 ACTGGTATCAACAGATC 592

RESULT 6
US-08-473-496-1/C
Sequence 1, Application US/08473496
Patent No. 5700660
GENERAL INFORMATION:
APPLICANT: Jack L. Leonard
APPLICANT: Peter E. Newburger
TITLE OF INVENTION: POSITIONAL CONTROL OF SELENIUM INSERTION
TITLE OF INVENTION: IN POLYPEPTIDES FOR X-RAY CRYSTALLOGRAPHY
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 MB
COMPUTER: IBM PS/2 Model 50Z or 55SX
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/473,496
FILING DATE: 07-JUN-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/277,492
FILING DATE: 19-JUL-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/066,680
FILING DATE: 24-MAY-1993
ATTORNEY/AGENT INFORMATION:
NAME: Fasse, J. Peter
REGISTRATION NUMBER: 32,983
REFERENCE/DOCKET NUMBER: 04020/078001

TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 832 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-473-496-1

Query Match 70.5% Score 14.8; DB 1; Length 832;
Best Local Similarity 88.9% Pred. No. 88;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcacagatc 20
|||||
DB 768 ACTGGATCACAGGACC 751

RESULT 7
US-08-454-028-3/c
Sequence 3, Application US/08454028
Patent No. 5849520
GENERAL INFORMATION:
APPLICANT: Jack L. Leonard
APPLICANT: Peter E. Newburger
TITLE OF INVENTION: POST-TRANSCRIPTIONAL GENE REGULATION BY
NUMBER OF SEQUENCES: 33
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM PS/2 Model 502 or 55SX
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/454,028
FILING DATE: May 30, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/066,680
FILING DATE: May 24, 1993
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Fasse, J. Peter
REGISTRATION NUMBER: 32,983
REFERENCE/DOCKET NUMBER: 04020/075001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 832 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-454-028-3

Query Match 70.5% Score 14.8; DB 2; Length 832;
Best Local Similarity 88.9% Pred. No. 88;

Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 3 actggatcacagatc 20
|||||
DB 768 ACTGGATCACAGGACC 751

RESULT 8
PCT-US94-05388-3/c
Sequence 3, Application PC/TUS9405388
GENERAL INFORMATION:
APPLICANT: Jack L. Leonard
APPLICANT: Peter E. Newburger
TITLE OF INVENTION: POST-TRANSCRIPTIONAL GENE
REGULATION BY TRACE
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM PS/2 Model 502 or 55SX
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/05388
FILING DATE: 16 May 1994
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/066,680
FILING DATE: May 24, 1993
ATTORNEY/AGENT INFORMATION:
NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,152
REFERENCE/DOCKET NUMBER: 04020/026001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 832
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
PCT-US94-05388-3

Query Match 70.5% Score 14.8; DB 5; Length 832;
Best Local Similarity 88.9% Pred. No. 88;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 3 actggatcacagatc 20
|||||
DB 768 ACTGGATCACAGGACC 751

RESULT 9
PCT-US95-09121-1/c
Sequence 1, Application PC/TUS9509121
GENERAL INFORMATION:
APPLICANT: University of Massachusetts
APPLICANT: Medical School
TITLE OF INVENTION: POSITIONAL CONTROL OF SILENIUM
TITLE OF INVENTION: INSERTION IN POLYPEPTIDES FOR
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.

STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM PS/2 Model 502 or
COMPUTER: 586X
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version
SOFTWARE: 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/09121
FILING DATE: 19 July 1995
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/277,492
FILING DATE: 19 July 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/473,496
FILING DATE: 7 June 1995
ATTORNEY/AGENT INFORMATION:
NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,162
REFERENCE/DOCKET NUMBER: 04020/078M01
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 832 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
PCT-US95-09121-1

Query Match 70.5%; Score 14.8; DB 5; Length 832;
Best Local Similarity 88.9%; Pred. No. 88;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcaacagatc 20
DB 768 ACTGGATCAACAGACC 751

RESULT 10
PCT-US96-07496-3/c
Sequence 3, Application PC/TUS9607496
GENERAL INFORMATION:
APPLICANT: University of Massachusetts Medical Center
TITLE OF INVENTION: POST-TRANSCRIPTIONAL GENE REGULATION BY
TITLE OF INVENTION: SELENIUM
NUMBER OF SEQUENCES: 33
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM PS/2 Model 502 or 586X
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US96/07496
FILING DATE:
CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/454,028
FILING DATE: May 30, 1995
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/066,680
FILING DATE: May 24, 1993
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Fasse, J. Peter
REGISTRATION NUMBER: 32,983
REFERENCE/DOCKET NUMBER: 04020/075M01
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 832 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
PCT-US96-07496-3

Query Match 70.5%; Score 14.8; DB 5; Length 832;
Best Local Similarity 88.9%; Pred. No. 88;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 actggatcaacagatc 20
DB 768 ACTGGATCAACAGACC 751

RESULT 11
US-08-698-551-13/c
Sequence 13, Application US/08698551
Patent No. 5712361
GENERAL INFORMATION:
APPLICANT: Lin, Lih-Ling
APPLICANT: Chen, Jennifer H.
APPLICANT: Schiavella, Andrea
APPLICANT: Graham, James
TITLE OF INVENTION: NOVEL TNF RECEPTOR DEATH DOMAIN LIGAND
TITLE OF INVENTION: PROTEINS AND INHIBITORS OF LIGAND BINDING
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 Cambridge Park Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: USA
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/698,551
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Brown, Scott A.
REGISTRATION NUMBER: 32,724
REFERENCE/DOCKET NUMBER: G15332D
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8224
TELEFAX: (617) 498-5851
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 3225 base pairs

TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 3..2846
US-08-698-551-13

Query Match 70.5%: Score 14.8; DB 1; Length 3225;
Best Local Similarity 88.9%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 ctggatcacagatct 21
||||| |||||||||
DB 504 CTGGTCACACAGATCT 487

RESULT 12
US-08-602-228-13/c
Sequence 13, Application US/08602228
Patent No. 5843675
GENERAL INFORMATION:
APPLICANT: Lin, Lih-Ling
APPLICANT: Chen, Jennifer H.
APPLICANT: Schievella, Andrea
TITLE OF INVENTION: NOVEL TNF RECEPTOR DEATH DOMAIN LIGAND
TITLE OF INVENTION: PROTEINS AND INHIBITORS OF LIGAND BINDING
NUMBER OF SEQUENCES: 16
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 Cambridgepark Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: USA
ZIP: 02140

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/602.228
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Brown, Scott A.
REGISTRATION NUMBER: 32,724
REFERENCE/DOCKET NUMBER: G15232C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8224
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 3225 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 3..2846
US-08-602-228-13

Query Match 70.5%: Score 14.8; DB 2; Length 3225;
Best Local Similarity 88.9%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 ctggatcacagatct 21
||||| |||||||||
DB 504 CTGGTCACACAGATCT 487

RESULT 13
US-08-533-901B-13/c
Sequence 13, Application US/08533901B
Patent No. 5852173
GENERAL INFORMATION:
APPLICANT: Lin, Lih-Ling
APPLICANT: Chen, Jennifer H.
APPLICANT: Schievella, Andrea
TITLE OF INVENTION: NOVEL TNF RECEPTOR DEATH DOMAIN LIGAND
TITLE OF INVENTION: PROTEINS AND INHIBITORS OF LIGAND BINDING
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 Cambridgepark Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: USA
ZIP: 02140

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/533.901B
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Brown, Scott A.
REGISTRATION NUMBER: 32,724
REFERENCE/DOCKET NUMBER: G15232
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8224
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 3225 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 3..2846
US-08-533-901B-13

Query Match 70.5%: Score 14.8; DB 2; Length 3225;
Best Local Similarity 88.9%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 ctggatcacagatct 21
||||| |||||||||
DB 504 CTGGTCACACAGATCT 487

RESULT 14
US-08-839-032A-13/c
Sequence 13, Application US/08839032A
Patent No. 5891675
GENERAL INFORMATION:
APPLICANT: Lin, Lih-Ling
APPLICANT: Chen, Jennifer H.
APPLICANT: Schievella, Andrea
TITLE OF INVENTION: NOVEL TNF RECEPTOR DEATH DOMAIN LIGAND PROTEINS

NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 CambridgePark Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: USA
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/839,032A
FILING DATE:
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Brown, Scott A,
REGISTRATION NUMBER: 32,724
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8224
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 3225 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 3..2846
US-08-839-032A-13
Query Match 70.5%; Score 14.8; DB 2; Length 3225;
Best Local Similarity 88.9%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
CY 4 ctgggatacaacagtatct 21
||||| |||||||||
DB 504 CTGGGTCCACAGTATCT 487
RESULT 15
US-08-839-031A-13/C
Sequence 13, Application US/08839031A
Patent No. 5948638
GENERAL INFORMATION:
APPLICANT: Lin, Lih-Ling
APPLICANT: Chen, Jennifer H.
APPLICANT: Schievella, Andrea
APPLICANT: Graham, James
TITLE OF INVENTION: NOVEL TNF RECEPTOR DEATH DOMAIN LIGAND
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 CambridgePark Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: USA
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/839,031A

FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Sprunger, Suzanne A.,
REGISTRATION NUMBER: 41,323
REFERENCE/DOCKET NUMBER: G15332BDIV
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8284
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 3225 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cDNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 3..2846
US-08-839-031A-13
Query Match 70.5%; Score 14.8; DB 2; Length 3225;
Best Local Similarity 88.9%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
CY 4 ctgggatacaacagtatct 21
||||| |||||||||
DB 504 CTGGGTCCACAGTATCT 487

Search completed: July 25, 2001, 05:18:52
Job time: 9215 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 25, 2001, 04:54:49 ; Search time 2762.24 Seconds
(without alignments)
71.866 Million cell updates/sec

Title: us-09-142-095-2

Perfect score: 21

Sequence: 1 ccactggatcacagatctc 21

Scoring table:

IDENTITY_NDC
Gapop 10.0 , Gapext 1.0

Searched: 1022815 seqs, 4726426750 residues

Total number of hits satisfying chosen parameters: 20456230

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	21	100.0	327	189	T71061	T71061 ycs0a04.r1
C 2	18.4	87.6	383	221	AA195421	AA195421 z336h09.s
C 3	18.4	87.6	1078	221	CNS04CBO	AA1284229 Telradon
C 4	17.8	84.8	297	169	BF764030	BF764030 IL2-CS004
C 5	17.8	84.8	425	3	AA195418	AA195418 z336g10.s
C 6	17.8	84.8	612	237	AA066229	AA066229 RPT-23-3
C 7	17.4	82.9	499	107	AU030401	AU030401 AU030401
C 8	17.4	82.9	499	107	AU030401	AU030401 AU030401
C 9	16.8	80.0	437	11	AA78238	AA78238 z197908.s
C 10	16.8	80.0	440	107	AU070697	AU070697 AU070697
C 11	16.8	80.0	450	107	AU094227	AU094227 AU094227
C 12	16.8	80.0	484	151	BP598564	BP598564 BP598564
C 13	16.8	80.0	520	155	BP557402	BP557402 BP557402
C 14	16.8	80.0	534	232	AO737479	AO737479 H8-3195.A
C 15	16.8	80.0	535	145	BF193631	BF193631 245281 MA
C 16	16.8	80.0	551	251	AE911013	AE911013 RPT-24-1
C 17	16.8	80.0	561	145	BF194355	BF194355 246337 MA
C 18	16.8	80.0	583	244	AZ483876	AZ483876 1M0310P03
C 19	16.8	80.0	628	6	AA392024	AA392024 LD10961.5
C 20	16.8	80.0	629	6	AA392019	AA392019 LD10961.5
C 21	16.8	80.0	698	107	AU094228	AU094228 AU094228
C 22	16.8	80.0	713	14	AA951996	AA951996 LD28992.5
C 23	16.8	80.0	881	150	BF578486	BF578486 602092908
C 24	16.4	78.1	306	160	BB520416	BB520416 BB520416
C 25	16.4	78.1	346	120	AW764293	AW764293 U74C08.X
C 26	16.4	78.1	522	142	BE942800	BE942800 EST422379
C 27	16.4	78.1	568	142	BE942802	BE942802 EST422381
C 28	16.4	78.1	683	240	AZ260166	AZ260166 RPT-23-1
C 29	16.4	78.1	694	219	AG013064	AG013064 Homo sapi
C 30	16.4	78.1	697	175	BC282828	BC282828 602405527
C 31	16.4	78.1	1055	146	BF304360	BF304360 601887268
C 32	16.4	78.1	1071	220	CNS02E0E	AA1193127 Tetradon
C 33	16.2	77.1	219	149	BF490633	BF490633 AT27168.5
C 34	16.2	77.1	306	131	BB336747	BB336747 BB336747
C 35	16.2	77.1	317	245	AZ504982	AZ504982 1M0345019
C 36	16.2	77.1	403	247	AZ647967	AZ647967 1M0314F17
C 37	16.2	77.1	412	118	AMS50028	AMS50028 hg30a04.x
C 38	16.2	77.1	416	172	BG019825	BG019825 dc68901.x
C 39	16.2	77.1	419	246	AZ604463	AZ604463 1M0425015
C 40	16.2	77.1	437	113	AM235361	AM235361 xm56a07.x
C 41	16.2	77.1	439	23	AI655576	AI655576 tt27n06.x
C 42	16.2	77.1	439	104	AI970357	AI970357 wg91c03.x
C 43	16.2	77.1	440	118	AMS50489	AMS50489 hg4se07.x
C 44	16.2	77.1	443	171	BF932393	BF932393 CM2-NT017
C 45	16.2	77.1	446	257	B98792	B98792 CIT-HSP-228

ALIGNMENTS

RESULT 1
T71061/c
LOCUS
DEFINITION
IMAGE:84078.5' similar to gb:M57899 UDP-GLUCURONOSYLTRANSFERASE 1A
PRECUSOR, MICROSOFT (HUMAN); mRNA sequence.
T71061
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS

327 bp mRNA
EST
01-MAR-1995
ycs0a04.r1 Stratagene liver (#937224) Homo sapiens cDNA clone
IMAGE:84078.5' similar to gb:M57899 UDP-GLUCURONOSYLTRANSFERASE 1A
PRECUSOR, MICROSOFT (HUMAN); mRNA sequence.
T71061
GI:685582
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 327)
Haller, L., Lennon, G., Becker, M., Bonaldo, M.F., Chippelli, B.,
Chissoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, N., Hawkins,
M., Hultman, M., Kucaba, T., Lacy, M., Le, N., Martin, E., Moore

FEATURES

FEATURES
SOURCE
1..327
/organism="Homo sapiens"
/db_xref="Gene:501135"
/db_xref="taxon:9606"
/clone="IMAGE:84078"
/clone_lib="Stratagene Liver (#937224)"
/sex="male"
/dev_stage="49 years old"
/lab_host="SOLR cells (kanamycin resistant)"
/note="Organ: liver; Vector: pBluescript SK; Site: 1: EcoRI
/ site: 2: XhoI; Cloned unidirectionally. Primer: Oligo
site: Hepatocytoma from normal male caucasian. Average insert
size: 1.1 kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5'
GATTGGGACGAG 3' -3' adaptor sequence: 5'
CTCGAGTTTCTTTTCTTTT 3'."
71 a 74 c 92 g 83 t 7 others

Query Match 100.0%; Score 21; DB 189; Length 327;
Best Local Similarity 100.0%; Pred. No. 2.2;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 2
AA195421
LOCUS
DEFINITION
IMAGE:84078.51 Soares, N.H.M.P., Si Homo sapiens cDNA clone IMAGE:665537 3'
sequence.
AA195421
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS

383 bp mRNA
EST
06-AUG-1997
z336h09.s1 Soares, N.H.M.P., Si Homo sapiens cDNA clone IMAGE:665537 3'
similar to contains element MSRI repetitive element; mRNA
sequence.
AA195421
GI:1785114
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 383)
Haller, L., Allen, M., Bowles, L., Dubuque, T., Gelsel, G., Jost, S.,
Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wyllie,
T., Waterston, R. and Wilson, R.
Wash-Merck EST Project 1997
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

JOURNAL

JOURNAL
COMMENT
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

FEATURES
SOURCE

Location/Qualifiers
1. .383

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/organism="Homo sapiens"
/cd_xref="CD8:5427569"
/db_xref="taxon:9606"
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/clone_1ib="Source:NbHPu_ST"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/ab_host="DH10B"
/mole="Organ: mixed (see below)" Vector: pT73D-Pac

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BASE COUNT	ORIGIN
108 a	83 c
72 g	112 t
8 others	

Query Match 87.6%; Score 18.4; DB 3; Length 383;
Best Local Similarity 90.5%; Prd. No. 49;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY      1 ccacgcggatcaacagtatct 21
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Db      180 CCACGCGCATCANCAAGTATCT 200

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RESULT	3
CNS04CB0/c	
LOCUS	
DEFINITION	CNS04CB0 1078 bp DNA
	tetraodon nigroviridis genome survey sequence PUC-Crl end of clone
	09NM1 of library G from tetraodon nigroviridis, genomic survey
	sequence.
ACCESSION	AF284229
VERSION	AL284229.1 GI:8022608
KEYWORDS	GSS; genome survey sequence.
SOURCE	tetraodon nigroviridis.
ORGANISM	tetraodon nigroviridis

REFERENCE
1 (bases 1 to 1078)
Roest-Crolius, H., Jallion, O., Dasilva, C., Frazar, C., Ribeiro, C.

TITLE	Characterization and repeat analysis of the compact genome of the freshwater purifierfish Tetraodon lineolus
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 1078)
ATTACHES	

TITLE	Human gene number estimate provided by genome wide analysis using Tetrahodon albigroviridis DNA sequence
JOURNAL	Unpublished
REFERENCE	3 (bases 1 to 1078)
AUTHORS	Gerosome

JOURNAL COMMENT
Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
This sequence is a single read and was generated as part of a large

scale clone-and sequencing project of the Tetradon nigroviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tetradon>.

Location/Qualifiers

1 1070

SOURCE

FEATURES

FEATURES
Source

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Query Match	87.6%;	Score 18.4;	DB 221;	Length 1078;
Best Local Similarity	95.08;	Pred. NO. 59;		
Matches 19;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0;
Qy 1 ccacttgatcacagcgtc 20				
Db 945 CCACGTGGATCACAGCTTC 926				

RESULT	4
BF764030/C	
LOCUS	
BEST	297 bp
DEFINITION	mRNA
ID	IL2-CSO048-301000-198-G10
ACCESSION	CSO048 Homo sapiens
VERSION	sapleins CDNA, mRNA sequence
KEYWORDS	BF764030.1 GI:12111930
SOURCE	EST.
	human.

ORGANISM	REFERENCE	AUTHORS
<i>Homo sapiens</i>		
Eularkyota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.		
1 (basses 1 to 237)		
Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Delgado, V.		

TITLE	Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE	20202663
COMMENT	Contact: Simpson A.J.G.

Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL:
<http://www.ludwig.org.br/scripts/gethtml2.pl?lcr-ll2&t2-ll2-CS0048>
301000-198-G10&t3-2008-10-30&t4-1)
Seq primer: puc 18 forward
High quality sequence step: 58.

FEATURES	Location/Qualifiers
source	1. .297

Location/Qualifiers
1. .297

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/note="Organ: colon_est; Vector: puc18; Site_1: Sma1;
Site_2: Sma1; A multi-library was made by cloning products
derived from ORS185 PCR (U.S. Letters Patent applications
No. 196,719 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

```


BASE COUNT 60 a 81 c 95 g 61 t
ORIGIN

Query Match 84.8% Score 17.8; DB 169; Length 297;
Best Local Similarity 90.5%; Pred. No. 1e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

0y 1 ccaactggatcaacagatct 21
|||||
Db 128 CCACGTGGAGCAGACGATCT 108

RESULT 5
AA195418 425 bp mRNA EST 06-AUG-1997
LOCUS z136g10.s1 Soares.NhMMPu.S1 Homo sapiens cDNA clone IMAGE:665538 3'
DEFINITION similar to contains element MERS repetitive element ; mRNA
SEQUENCE.
AA195418
VERSION AA195418.1 GI:1785111
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 425)
Hiller, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
Kucab, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B.,
Schellenberg, K., Stepien, M., Tan, F., Theising, B., White, Y., Wyllie,
'T., Waterston, R., and Wilson, R.
Mashu-Merck EST Project 1997
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8500, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.edu
This clone is available royalty-free through LIND; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1873 Std Error: 0.00
High quality sequence stop: 337.
Location/Qualifiers
1..425
/organism="Homo sapiens"
/db_xref="GDB:5427570"
/db_xref="taxon:9606"
/clone="IMAGE:665538"
/clone_id="Soares.NhMMPu.S1"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/note="Organ: mixed (see below); Vector: pT7T3D-Pac
(Pharmacia) with a modified polylinker; Site 1: Not I;
Site 2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NDHM, pregnant uterus
NBHPU, and fetal heart NBH129) were mixed, and ss circles
were made in vitro. Following BAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of 1 M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."

BASE COUNT 129 a 85 c 80 g 129 t
ORIGIN

Query Match 84.8% Score 17.8; DB 3; Length 425;
Best Local Similarity 90.5%; Pred. No. 1e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
0y 1 ccaactggatcaacagatct 21

Db 180 CCACGTGGATCAGACGATCT 200
|||||

RESULT 6
A2066229/c 612 bp DNA GSS 30-MAR-2000
LOCUS RPCI-23-392B4.TV RPCI-23 Mus musculus genomic clone RPCI-23-392B4,
DEFINITION DNA sequence.
A2066229
ACCESSION A2066229.1 GI:7357481
VERSION
KEYWORDS
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 612)
Zhao, S., Nierman, W., Feldblyum, T., Malek, J., Shatsman, S., Akintet,
'B., Levins, M., McGann, S., Tsegaye, G., Geer, K., Krol, M., de Jong, P.,
and Fraser, C.M.
Mouse BAC End Sequences from Library RPCI-23
Unpublished (1998)
Contact: Shuying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the mouse BAC library RPCI-23. For BAC
library availability, please contact Pieter de Jong
(pieter@edj.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/orderingframe.htm>)
or from Resea ch Genetics (info@resgen.com). BAC end page:
http://www.tigr.org/tldb/bac_ends/mouse/Bac_end_intro.html
Plate: 392 Row: B Column: 4
Seq primer: 75
Class: BAC ends.
Location/Qualifiers
1..612
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="RPCI-23-392B4"
/clone_id="RPCI-23"
/sex="Female"
/lab_host="DH10B"
/note="Organ: Kidney/Brain; Vector: pBAC3.6; Site 1:
EcoRI; Site 2: EcoRI; Female C57BL/6J mouse kidney and/or
brain genomic DNA was isolated and partially digested
with a combination of EcoRI and EcoRI Methylase. Size
selected DNA was cloned into the pBAC3.6 vector at the
EcoRI sites. The ligation products were transformed into
DH10B electrocompetent cells (BRL Life Technologies)."

BASE COUNT 176 a 121 c 116 g 199 t
ORIGIN

Query Match 84.8% Score 17.8; DB 237; Length 612;
Best Local Similarity 90.5%; Pred. No. 1.1e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
0y 1 ccaactggatcaacagatct 21
|||||
Db 540 CCCCTGGATCAGACGATCT 520

RESULT 7
AU030401/c 499 bp mRNA EST 19-OCT-1998
LOCUS AU030401 Rice cDNA from immature leaf including apical meristem
DEFINITION Oryza sativa cDNA clone EST014748, mRNA sequence.
ACCESSION AU030401

This is a subtracted version of the original Soares fetal

liver spleen INFLS library. 1st strand cDNA was primed with a Pac I - oligo(dT) primer [5', AACTGGAAGAATTAATTAAGATCTTTTTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Pac I and cloned into the Pac I and Eco RI sites of the modified p773 vector. Library constructed through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 116 a 104 c 99 g 118 t
ORIGIN

Query Match 80.0%; Score 16.8; DB 11; Length 437;
Best Local Similarity 90.0%; Pred. NO. 3.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 cactgggatcaacagtatct 21
|||||
Db 315 CACTGGGATCAACAAATATTT 296

RESULT 10
AU070697/c 440 bp mRNA EST 10-JUN-1999
LOCUS
DEFINITION AU070697 Rice cDNA from young root Oryza sativa cDNA clone
R10115_1A, mRNA sequence.
ACCESSION AU070697
VERSION AU070697
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.

REFERENCE
AUTHORS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
TITLE Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
JOURNAL Ehrhartoidae; Oryzeae; Oryza.
COMMENT 1 (bases 1 to 440)
Rice cDNA from young root
Yamamoto, K. and Sasaki, T.
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program
2-1-2 Kannondai, Tsukuba
Ibaraki,
Japan 305
Tel: 0298-38-7441
Fax: 0298-38-7468
Email: tsasaki@abr.affrc.go.jp
PROJECT "RGP".

FEATURES
Source
Location/Qualifiers
1..440
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="R10115_1A"
/clone_lib="Rice cDNA from young root"
/tissue_type="young root"

BASE COUNT 126 a 86 c 94 g 132 t
ORIGIN

Query Match 80.0%; Score 16.8; DB 107; Length 440;
Best Local Similarity 90.0%; Pred. NO. 3.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 cactgggatcaacagtatct 21
|||||
Db 29 CACTGGGATCAGCAGTATCT 10

RESULT 11
AU094227/c 450 bp mRNA EST 30-JUN-2000
LOCUS
DEFINITION AU094227 Rice panicle at flowering stage Oryza sativa cDNA clone

ACCESSION E3685, mRNA sequence.
VERSION AU094227
KEYWORDS AU094227.1 GI:8856909
EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.

REFERENCE
AUTHORS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
TITLE Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
JOURNAL Ehrhartoidae; Oryzeae; Oryza.
COMMENT 1 (bases 1 to 450)
Rice cDNA from panicle at flowering stage (2000)
Unpublished (2000)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program
2-1-2 Kannondai, Tsukuba
Ibaraki,
Japan 305
Tel: 0298-38-7441
Fax: 0298-38-7468
Email: tsasaki@abr.affrc.go.jp
PROJECT "RGP".

FEATURES
source
Location/Qualifiers
1..450
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="E3685"
/clone_lib="Rice panicle at flowering stage"
/dev_stage="flowering stage"
/note="Organ: panicle; Rice cDNA from panicle at flowering stage"

BASE COUNT 117 a 91 c 121 g 120 t 1 others
ORIGIN

Query Match 80.0%; Score 16.8; DB 107; Length 450;
Best Local Similarity 90.0%; Pred. NO. 3.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 cactgggatcaacagtatct 21
|||||
Db 254 CACTGGGATCAGCAGTATCT 235

RESULT 12
BF598564/c 484 bp mRNA EST 12-DEC-2000
LOCUS
DEFINITION BF598564 Glycine max cDNA clone GENOME SYSTEMS CLONE ID:
sv19a09.y1 Gm-cl057 similar to SW:PLAS_LYC5S P17340 PLASTOCYANIN
Gm-cl057-17 5', similar to SW:PLAS_LYC5S P17340 PLASTOCYANIN
PRECURSOR. [1]; mRNA sequence.

ACCESSION BF598564
VERSION BF598564.1 GI:11690888
KEYWORDS EST.
SOURCE soybean.
ORGANISM Glycine max

REFERENCE
AUTHORS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
TITLE Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
JOURNAL Rosidae; eurosids 1; Fabales; Fabaceae; Papilionoideae; Glycine.
COMMENT 1 (bases 1 to 484)
Shoemaker, R., Kelm, P., Vodkin, L., Erpelting, J., Coryell, V., Khanna
A., Bolla, B., Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C.,
Wyllie, T., Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers
Y., Person, B., Swaller, F., Gibbons, M., Pape, D., Harvey, N., Schurk
R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann
R., Waterston, R. and Wilson, R.
Public Soybean EST Project
Unpublished (1999)
Contact: Shoemaker R/Public Soybean EST Project
Public Soybean EST Project
Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu

This clone is available through: Genome Systems, Inc. 4633 World Parkway Circle St. Louis, Missouri 63134 for further information call: (800) 430-0030 or (314) 427-3222 FAX: (888) 919-3324 or (314) 427-3324 or contact: clones@genomesystems.com or info@genomesystems.com web site: www.genomesystems.com
Trace considered overall poor quality
High quality sequence stop: 1.

FEATURES

source

Location/Qualifiers

1..484

/organism="Glycine max"

/db.xref="taxon:3847"

/clone="GENOME SYSTEMS CLONE ID: Gm-cl057-17"

/tissue_type="degenerating cotyledons, 2 week old seedling"

/lab_host="DH10B"

/note="Vector: Bluescript II SK⁺ Site.1: EcoRI; Site.2: XhoI; The cDNA library was constructed from mRNA isolated from degenerating cotyledons of 2 week old seedlings from p1468916. Complementary DNA was synthesized from mRNA using a primer consisting of a poly(dT) sequence with a XhoI restriction site. EcoRI adapters were ligated to the blunt-ended cDNA fragments followed by XhoI digestion. The cDNA fragments were directionally cloned into the EcoRI-XhoI restriction site of the Bluescript vector. The ligated cDNA fragments were transformed into DH10B host cells (GibcoBRL). This library was constructed in the laboratory of Dr. Randy Shoemaker."

BASE COUNT 117 a 133 c 121 g 113 t

ORIGIN

Query Match 80.0%; Score 16.8; DB 151; Length 484;
Best Local Similarity 90.0%; Pred. No. 3.4e+02;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Y 1 ccactgggatacagatc 20

DB 435 CCATGGCATCAACAGTATC 416

RESULT 13

LOCUS

BG557402 520 bp mRNA EST 10-APR-2001

DEFINITION

EML 43 E08.b1.A002 Embryo 1 (EML) Sorghum bicolor cDNA, mRNA

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BG557402 520 bp mRNA EST 10-APR-2001
EML 43 E08.b1.A002 Embryo 1 (EML) Sorghum bicolor cDNA, mRNA
sequence.
BG557402
BG557402.1 GI:13586400
EST.
Sorghum.
Sorghum bicolor
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
1 (bases 1 to 520)
Reid,S.P., Cordonier-Pratt,M.-M., Gingie,A. and Pratt,L.H.
An EST database from Sorghum: developing embryos
Unpublished (2000)
Contact: Cordonier-Pratt MM
Department of Botany
The University of Georgia
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 542 1805
Email: mpratt@uga.edu
Sequences have been trimmed to exclude PolyA, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.
Seq primer: JEN REV

FEATURES

source

Location/Qualifiers

1..520

/organism="Sorghum bicolor"

/db.xref="taxon:4558"

/clone.lib="Embryo 1 (EML)"

/note="Organ: Embryos germinated for 24 hr; Vector: Bluescript II from Lambda Zap II; Site.1: XhoI; Site.2: EcoRI; The library was made from poly-A RNA in the cloning vector lambda Zap II. Clones to be sequenced were prepared by mass excision."

BASE COUNT 121 a 164 c 112 g 122 t 1 others

ORIGIN

Query Match 80.0%; Score 16.8; DB 155; Length 520;
Best Local Similarity 90.0%; Pred. No. 3.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Y 2 cactgggatacagatc 21

DB 2 CACGAGATCAACAGTACT 21

RESULT 14

LOCUS

AO737479 534 bp DNA GSS 16-JUL-1999

HS_3195_A2_B02_T7C CIR Approved Human Genomic Sperm Library D Homo

sapiens genomic clone Plate=3195 Col=4 Row=C, DNA sequence.

ACCESSION AO737479.1 GI:5515001

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 534)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,D., Zhao,S., Adams,M.D. and
Hood,L.

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)

TITLE

JOURNAL

MEDLINE

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones may be purchased from Research Genetics (info@resgen.com).
BAC end Web Server: http://www.htsc.washington.edu
Plate: 3195 Row: C Column: 4
Seq primer: T7
Class: BAC ends

High quality sequence stop: 534.

FEATURES

source

Location/Qualifiers

1..534

/organism="Homo sapiens"

/db.xref="taxon:9606"

/clone.lib="Plate=3195 Col=4 Row=C"

/sex="male"

/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in E-Coli DH10B"

ORIGIN

BASE COUNT 140 a 120 c 121 g 146 t 7 others

Query Match 80.0%; Score 16.8; DB 232; Length 534;
Best Local Similarity 90.0%; Pred. No. 3.5e+02;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 cactggatcacagatct 21
 ||||| ||||| ||
 Db 210 CACTGGAGCAGACTACT 191

RESULT 15

BF193631 535 bp mRNA EST 02-NOV-2000
 LOCUS 245281 MARC 2Pig sus scrofa cDNA 5', mRNA sequence.
 DEFINITION
 ACCESSION BF193631
 VERSION BF193631.1 GI:11077000
 KEYWORDS EST.
 SOURCE
 ORGANISM
 Pig.
 Sus scrofa
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.

REFERENCE

1 (bases 1 to 535)
 Fahrenkrug, S.C., Freking, B.A., Rohrer, G.A., Smith, T.P.L., Casas, E.,
 Stone, R.T., Heaton, M.P., Grose, W.M., Bennett, G.A., Laegreid, W.W.
 and Keele, J.W.
 Design and use of two pooled tissue normalized cDNA libraries for
 EST discovery in swine
 Unpublished (2000)

TITLE

JOURNAL

COMMENT
 Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smithemall.marc.usda.gov
 Single pass sequencing. Bases called and alt-trimmed with phred
 v0.980904.e. Vector identified by cross-match with the -mismatch 18
 and -mismatch 12 options.
 PCR Primers
 FORWARD: AGGAACAGTATGACAT
 BACKWARD: GTTTCAGTACAGAC
 Plate: 74 row: K column: 12
 Seq primer: ATTAGTGACACTATAG.
 Location/Qualifiers

FEATURES

source
 1..535
 /organism="Sus scrofa"
 /db_xref="taxon:9823"
 /clone_lib="MARC 2Pig"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /note="Vector: pCMV SPORT6; Site 1: XbaI; Site 2: XhoI;
 Library made from pooled tissue from testis, ovary,
 endometrium, hypothalamus, pituitary, and placenta."
 BASE COUNT 163 a 136 c 118 g 118 t
 ORIGIN

Query Match 80.0%; Score 16.8; DB 145; Length 535;
 Best local similarity 90.0%; Pred. No. 3.5e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ccactggatcacagatc 20
 ||||| ||||| ||
 Db 514 CCACTGGATCAGACTACT 533

Search completed: July 25, 2001, 04:54:51
 Job time: 10424 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OW nucleic - nucleic search, using sw model

Run on: July 25, 2001, 05:16:58 ; Search time 1290.33 Seconds

(without alignments)
239.749 Million cell updates/sec

Title: US-09-142-095-3

Perfect score: 20
Sequence: 1 gtacagtgcacagtcacac 20

Scoring table: IDENTITY_NTC
Gapop 10.0 , Gapext 1.0

Searched: 1344157 segs, 7733874588 residues

Total number of hits satisfying chosen parameters: 2688314

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl:*

- 1: gb_pat1:*
- 2: gb_da2:*
- 3: gb_da3:*
- 4: gb_in1:*
- 5: gb_in2:*
- 6: gb_in3:*
- 7: gb_cm:*
- 8: gb_ov:*
- 9: gb_pat2:*
- 10: gb_pat3:*
- 11: gb_ph:*
- 12: gb_p11:*
- 13: gb_p12:*
- 14: gb_p13:*
- 15: gb_p14:*
- 16: em_da1:*
- 17: em_da2:*
- 18: em_fun:*
- 19: em_hrgo_hum:*
- 20: em_hrgo_inv:*
- 21: em_hrgo_rtd:*
- 22: em_hrg_hum1:*
- 23: em_hrg_hum2:*
- 24: em_hrg_hum3:*
- 25: em_hrg_hum4:*
- 26: em_hrg_hum5:*
- 27: em_hrg_hum6:*
- 28: em_hrg_hum7:*
- 29: em_hrg_hum8:*
- 30: em_hrg_inv1:*
- 31: em_hrg_inv2:*
- 32: em_hrg_other:*
- 33: em_hrg_rtd:*
- 34: em_hum1:*
- 35: em_hum2:*
- 36: em_hum3:*
- 37: em_hum4:*
- 38: em_hum5:*
- 39: em_hum6:*
- 40: em_hum7:*
- 41: em_in:*
- 42: em_cm:*
- 43: em_or:*

44: em_ov:*

- 45: em_pat:*
- 46: em_ph:*
- 47: em_p1:*
- 48: em_rtd:*
- 49: em_sl:*
- 50: em_sy:*
- 51: em_un:*
- 52: em_v1:*
- 53: gb_stg1:*
- 54: gb_stg2:*
- 55: gb_stg3:*
- 56: gb_sy:*
- 57: gb_un:*
- 58: gb_v11:*
- 59: gb_v12:*
- 60: gb_hrg1:*
- 61: gb_hrg2:*
- 62: gb_hrg3:*
- 63: gb_hrg4:*
- 64: gb_hrg5:*
- 65: gb_hrg6:*
- 66: gb_hrg7:*
- 67: gb_hrg8:*
- 68: gb_hrg9:*
- 69: gb_hrg10:*
- 70: gb_hrg11:*
- 71: gb_hrg12:*
- 72: gb_hrg13:*
- 73: gb_hrg14:*
- 74: gb_hrg15:*
- 75: gb_hrg16:*
- 76: gb_hrg17:*
- 77: gb_hrg18:*
- 78: gb_hrg19:*
- 79: gb_hrg20:*
- 80: gb_hrg21:*
- 81: gb_hrg22:*
- 82: gb_hrg23:*
- 83: gb_hrg24:*
- 84: gb_hrg25:*
- 85: gb_p1:*
- 86: gb_p2:*
- 87: gb_p3:*
- 88: gb_p4:*
- 89: gb_p5:*
- 90: gb_p6:*
- 91: gb_p7:*
- 92: gb_p8:*
- 93: gb_p9:*
- 94: gb_rtd:*
- 95: gb_rtd:*
- 96: gb_in4:*
- 97: gb_p10:*
- 98: em_da3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	20	100.0	20 9 A65502	A65502 Sequence 3
2	20	100.0	531 89 AF352795	AF352795 Homo sapi
3	20	100.0	541 89 AF180372	AF180372 Homo sapi
4	20	100.0	620 9 A65504	A65504 Sequence 5
5	20	100.0	3341 91 D87674	D87674 Homo sapien
6	20	100.0	68770 86 AC006985	AC006985 Homo sapi
7	20	100.0	116619 70 AC026497	AC026497 Homo sapi
8	20	100.0	119872 89 AF297093	AF297093 Homo sapi


```

AF180372      541 bp      DNA      PRI      05-OCT-1995
LOCUS      AF180372      Homo sapiens bilirubin UDP-glucuronosyltransferase 1-1 (UGT1) gene,
DEFINITION      UGT1*1 allele, partial cds.
ACCESSION      AF180372
VERSION      AF180372.1      GI:6010649
KEYWORDS
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE      1 (bases 1 to 541)
AUTHORS      Kutlar,F., Stromek,E., Leitner,C., Nechtman,J. and Kutlar,A.
TITLE      Deletion of the TATA box polymorphism of the human bilirubin
JOURNAL      UDP-glucuronosyltransferase 1-1 gene (UGT1*1) in a patient with
              sickle cell anemia
              Unpublished
REFERENCE      2 (bases 1 to 541)
AUTHORS      Kutlar,F., Stromek,E., Leitner,C., Nechtman,J. and Kutlar,A.
TITLE      Direct Submission
JOURNAL      Submitted (24-AUG-1999) Medicine, Hematology/Oncology-Sickle Cell
              Center, Medical College of Georgia, 15th Street, AC-1000, Augusta,
              GA 30912, USA
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Best Local Similarity 100.0%; Pred. No. 0.58;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1      gtacagtgcacagtcacac 20
DB      228      gtacagtgcacagtcacac 247

RESULT      4
LOCUS      A65504
DEFINITION      A65504      620 bp      DNA      PAT      29-MAR-1999
              Sequence 5 from Patent WO9732042.
ACCESSION      A65504
VERSION      A65504.1      GI:4531239
KEYWORDS
SOURCE      unidentified.
ORGANISM      unidentified.
REFERENCE      1 (bases 1 to 620)
AUTHORS      Birrell,B.
TITLE      DRUG TRIAL ASSAY SYSTEM
JOURNAL      Patent: WO 9732042-A 5 04-SEP-1997;
              UNIV DUNDEE (GB)
              Other publication AU 2224197 19970916.
COMMENT      location/Qualifiers
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1      gtacagtgcacagtcacac 20
DB      508      gtacagtgcacagtcacac 527

RESULT      5
LOCUS      D87674
DEFINITION      D87674      3341 bp      DNA      PRI      14-APR-2000
              Homo sapiens gene for bilirubin UDP-glucuronosyltransferase 1,
              promoter region and partial cds.
ACCESSION      D87674
VERSION      D87674.1      GI:3059176
KEYWORDS      bilirubin UDP-glucuronosyltransferase 1.
SOURCE      Homo sapiens DNA.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE      1 (sites)
AUTHORS      Ueyama,H., Koike,O., Soeda,Y., Sato,H., Satoh,Y., Okubo,I. and
              Doi,Y.
TITLE      Analysis of the promoter of human bilirubin
              UDP-glucuronosyltransferase gene (UGT1*1) in relevance to Gilbert's
              syndrome
JOURNAL      Hepatol. Res. 9, 152-163 (1997)
REFERENCE      2 (bases 1 to 3341)
AUTHORS      Ueyama,H.
TITLE      Direct Submission
JOURNAL      Submitted (04-SEP-1996) to the DDBJ/EMBL/GenBank databases. Hisao
              Ueyama, Shiga University of Medical Science, Department of Medical
              Biochemistry, Seto, Otsu, Shiga 520-21, Japan (Tel:077-548-2162,
              Fax:077-548-2164)
COMMENT      Sequence updated (08-Jan-1997) by: Hisao Ueyama.
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BASE COUNT 893 a 695 c 803 g 950 t
ORIGIN

Query Match 100.0%; Score 20; DB 91; Length 3341;
Best Local Similarity 100.0%; Pred. No. 0.35;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gtcacgtgacacagtcacac 20
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Db 3088 gtcacgtgacacagtcacac 3107

RESULT 6
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LOCUS
DEFINITION Homo sapiens BAC clone RP11-154L24 from 2, complete sequence.
AC006985
VERSION AC006985.2 GI:5732165
KEYWORDS
HTG.
SOURCE
human.

ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Sulston, J.E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
REFERENCE
AUTHORS Gattung, S., Stoneking, T. and Davidson, T.
TITLE The sequence of Homo sapiens BAC clone RP11-154L24
JOURNAL Unpublished
3 (bases 1 to 68770)
REFERENCE
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (05-MAR-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 68770)
REFERENCE
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (13-AUG-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 68770)
REFERENCE
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (22-OCT-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
6 (bases 1 to 68770)
REFERENCE
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (21-DEC-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Aug 13, 1999 this sequence version replaced g1:4337236.

COMMENT
Center: Washington University Genome Sequencing Center
Center code: WUGSC

Web site: <http://genome.wustl.edu/gsc>
Contact: saplens@wustl.edu
Summary Statistics
Center project name: H_NH0154L24

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RP11-154L24 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Moon, F.Y., Zhao, B., Firengen, E., Tatem, M., Catalanese, J.J. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)
VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-332L11, 200 bp overlap. Actual start of this clone is at base position 85134 of RP11-332L11; actual end is at base position 68770 of RP11-154L24.

The clone RP11-154L24 contains a tandem repeat from base positions 38234 to 39039, this region contains some low quality data. The assembly is consistent with the restriction digest information.

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45619	48728	:contigs of 3110 bp in length
48729	48828	:gap of 100 bp in length
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52100	52199	:gap of 100 bp in length
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56748	56877	:gap of 100 bp in length
56848	60467	:contigs of 3620 bp in length
60468	60577	:gap of 100 bp in length
60568	64947	:contigs of 4380 bp in length
64948	65047	:gap of 100 bp in length
65048	69431	:contigs of 4384 bp in length
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69532	74167	:contigs of 4636 bp in length
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* 100588.100687: gap of 100 bp
* 100588.105884: contig of 5197 bp in length
* 105885.105984: gap of 100 bp
* 105885.111486: contig of 5502 bp in length
* 111487.111586: gap of 100 bp
* 111887.117881: contig of 6195 bp in length
* 117882.125343: contig of 7462 bp in length
* 125344.125443: gap of 100 bp
* 125444.135058: contig of 7055 bp in length
* 132509.122608: gap of 100 bp
* 132609.141180: contig of 8582 bp in length
* 141191.141290: gap of 100 bp
* 141191.155556: contig of 11266 bp in length
* 152557.152656: gap of 100 bp
* 152857.163590: contig of 10934 bp in length
* 163591.163680: gap of 100 bp
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 Best local similarity 100.0%; Pred. No. 0.48;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gtccagtcacacagtcacac 20
 Db 174939 gtccagtcacacagtcacac 174938
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 DEFINITION Pongo pygmaeus UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
 promoter region and partial cds.
 ACCESSION AF135466
 VERSION AF135466.1 GI:6456549
 KEYWORDS
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 ORGANISM
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 Pongo pygmaeus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Pongo.
 Hall, D., Ybazeza, G., Destro-Bisoli, G., Petzi-Erler, M.L. and Di
 Rienzo, A.
 Variability at the uridine diphosphate glucuronosyltransferase 1A1
 promoter in human populations and primates
 Pharmacogenetics (1999) in press
 2 (bases 1 to 200)
 Ybazeza, G., Hall, D. and Di Rienzo, A.
 Direct Substitution
 Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924
 E. 57th street, Chicago, IL 60637, USA
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 DEFINITION Trachypithecus obscurus UDP-glucuronosyltransferase 1A1 (UGT1A1)
 gene, promoter region and partial cds.
 ACCESSION AF135467
 VERSION AF135467.1 GI:6456551
 KEYWORDS
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 ORGANISM
 dusky leaf monkey.
 Trachypithecus obscurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
 Colobinae; Trachypithecus.
 Hall, D., Ybazeza, G., Destro-Bisoli, G., Petzi-Erler, M.L. and Di

TITLE
Rienzo, A.
Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates

JOURNAL
Pharmacogenetics (1999) In press

REFERENCE
2 (bases 1 to 208)

AUTHORS
Ybazaeta, G., Hall, D. and Di Rienzo, A.

TITLE
Direct Submission

JOURNAL
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924 E. 57th Street, Chicago, IL 60637, USA

FEATURES
SOURCE
Location/Qualifiers
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Query Match
Best Local Similarity 92.0%; Score 18.4; DB 89; Length 208;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gtcacgtgacacagtcacaac 20
|||||

DB 30 gtcacgtgacacagtcacaac 49

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AF135462 234 bp DNA PRI 21-NOV-1999
LOCUS
DEFINITION Pan paniscus UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
promoter region and partial cds.
ACCESSION
AF135462
AF135462.1 GI:6456541
KEYWORDS
pygmy chimpanzee.
SOURCE
Pan paniscus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.
1 (bases 1 to 234)
Hall, D., Ybazaeta, G., Destro-Bisoli, G., Petzl-Erler, M.L. and Di
Rienzo, A.
Variability at the uridine diphosphate glucuronosyltransferase 1A1
promoter in human populations and primates
Pharmacogenetics (1999) In press
2 (bases 1 to 234)
Ybazaeta, G., Hall, D. and Di Rienzo, A.
Direct Submission
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924
E. 57th Street, Chicago, IL 60637, USA

FEATURES
SOURCE
Location/Qualifiers
1..234
/organism="Pan paniscus"
/db_xref="taxon:9597"
/gene="UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
/db_xref="GI:6456544"
/protein_id="AA09174.1"
/translation="MAVESODRPLVGLLCLVLPVCHAG"

BASE COUNT
50 a 53 c 69 g 62 t

ORIGIN

Query Match
Best Local Similarity 92.0%; Score 18.4; DB 89; Length 234;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gtcacgtgacacagtcacaac 20
|||||

DB 46 gtcacgtgacacagtcacaac 65

RESULT 13
MUSPC326A 4201 bp mRNA ROD 27-APR-1999
LOCUS
DEFINITION Mus musculus protein P0326 mRNA, complete cds.

BASE COUNT
50 a 53 c 69 g 62 t

ORIGIN

Query Match
Best Local Similarity 92.0%; Score 18.4; DB 89; Length 234;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gtcacgtgacacagtcacaac 20
|||||

DB 46 gtcacgtgacacagtcacaac 65

RESULT 12
AF135463 234 bp DNA PRI 21-NOV-1999
LOCUS
DEFINITION Pan troglodytes UDP-glucuronosyltransferase 1A1 (UGT1A1) gene,
promoter region and partial cds.
ACCESSION
AF135463
AF135463.1 GI:6456543
KEYWORDS
chimpanzee.
SOURCE
Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.
1 (bases 1 to 234)
Hall, D., Ybazaeta, G., Destro-Bisoli, G., Petzl-Erler, M.L. and Di
Rienzo, A.
Variability at the uridine diphosphate glucuronosyltransferase 1A1
promoter in human populations and primates
Pharmacogenetics (1999) In press
2 (bases 1 to 234)
Ybazaeta, G., Hall, D. and Di Rienzo, A.
Direct Submission
Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924
E. 57th Street, Chicago, IL 60637, USA

FEATURES
SOURCE
Location/Qualifiers
1..234
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/gene="UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
/db_xref="GI:6456544"
/protein_id="AA09174.1"
/translation="MAVESODRPLVGLLCLVLPVCHAG"

BASE COUNT
50 a 53 c 69 g 62 t

ORIGIN

Query Match
Best Local Similarity 92.0%; Score 18.4; DB 89; Length 234;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gtcacgtgacacagtcacaac 20
|||||

DB 46 gtcacgtgacacagtcacaac 65

RESULT 13
MUSPC326A 4201 bp mRNA ROD 27-APR-1999
LOCUS
DEFINITION Mus musculus protein P0326 mRNA, complete cds.

Query Match	87.08;	Score 17.4;	DB 94;	Length 4201;
Best Local Similarity	94.78;	Pred. No. 20;		
Matches 18;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0
QY	1 gtcacgtacacagctcaaa 19			

REFERENCE	AUTHORS	TITLE
1 (bases 1 to 207)	Hall, D., Ybazeta, G., Destro-Bisol, G., Petzl-Erler, M. L. and Di Rienzo, A.	Variability at the uridine diphosphate glucuronosyltransferase 1A1 promoter in human populations and primates

JOURNAL Pharmacogenetics (1999) In press
REFERENCE 2 (bases 1 to 207)
AUTHORS Ybazaeta,G., Hall,D. and Di Rienzo,A.
TITLE Direct Submission
JOURNAL Submitted (18-MAR-1999) Human Genetics, University of Chicago, 924
E. 57th Street, Chicago, IL 60637, USA
FEATURES
SOURCE location/Qualifiers
1..207

organism="Cebus apella"
/db_xref="taxon:9515"
<144..>207

gene mRNA

gene "UGT1A1"
/product="UDP-glucuronosyltransferase 1A1"
<144..>207

CDS "UGT1A1"
144..>207

/codon_start=1
/product="UDP-glucuronosyltransferase 1A1"
/protein_id="AA09181.1"

/db_xref="GI:6456558"
/translation="MPACGPAVACAGPGSVPCWE"
BASE COUNT 43 a 46 c 62 g 56 t

ORIGIN

Query Match 84.08; Score 16.8; DB 89; Length 207;
Best Local Similarity 90.08; Pred. No. 52;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 gtacgtgacacagtcacac 20
|||||

Db 23 gtctctgacacatcattac 42
|||||

Search completed: July 25, 2001, 05:17:09
Job time: 9232 sec